

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification 6: C07K 16/00, C07H 21/04, C12P 21/06, A61K 38/00</p>	<p>A1</p>	<p>(11) International Publication Number: WO 99/26980 (43) International Publication Date: 3 June 1999 (03.06.99)</p>
<p>(21) International Application Number: PCT/US98/24976 (22) International Filing Date: 19 November 1998 (19.11.98) (30) Priority Data: 08/976,096 21 November 1997 (21.11.97) US 60/099,160 4 September 1998 (04.09.98) US (63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Applications US 08/976,096 (CIP) Filed on 21 November 1997 (21.11.97) US 60/099,160 (CIP) Filed on 4 September 1998 (04.09.98) (71) Applicant (for all designated States except US): BETH ISRAEL DEACONESS MEDICAL CENTER [US/US]; 330 Brookline Avenue, Boston, MA 02215 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): TERHORST, Cornelis, P. [US/US]; 72 Fayerweather Street, Cambridge, MA 02215 (US). SAYOS-ORTEGA, Joan [ES/US]; 143 Park Drive #12, Boston, MA 02110-2214 (US).</p>		<p>(74) Agent: BIEKER-BRADY, Kristina; Clark &amp; Elbing LLP, 176 Federal Street, Boston, MA 02110-2214 (US). (81) Designated States: AU, CA, JP, NZ, US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  Published With international search report.</p> <p>9E 51</p>
<p>(54) Title: METHODS AND REAGENTS FOR THE UTILIZATION OF SAP FAMILY MEMBER PROTEINS, NOVEL SIGNAL TRANSDUCTION REGULATORS (57) Abstract  The invention provides novel SAP proteins and nucleic acid sequences. The invention also identifies the SAP family, members of which are novel signal transduction regulators. Also provided are SAP family polypeptides, SAP family member polypeptide specific antibodies, and methods for modulating SH2 domain-containing protein-mediated signal transduction, in particular, antigen-specific T cell activation. In addition, methods are provided for detecting compounds for modulating SH2 domain-containing protein-mediated signal transduction.</p>		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon	KR	Republic of Korea	PL	Poland		
CN	China	KZ	Kazakhstan	PT	Portugal		
CU	Cuba	LC	Saint Lucia	RO	Romania		
CZ	Czech Republic	LI	Liechtenstein	RU	Russian Federation		
DE	Germany	LK	Sri Lanka	SD	Sudan		
DK	Denmark	LR	Liberia	SE	Sweden		
EE	Estonia			SG	Singapore		

METHODS AND REAGENTS FOR THE UTILIZATION OF SAP FAMILY  
MEMBER PROTEINS. NOVEL SIGNAL TRANSDUCTION REGULATORS

5

Background of the Invention

The invention relates to modulation of SH2 domain-containing protein-mediated signal transduction generally, and in particular antigen-specific T cell activation.

The signal transduction cascade emanating from a cell surface-localized  
10 receptor into the nucleus of the cell where activation-specific genes are upregulated involves a very large number of molecules, including kinases, phosphatases, and adaptor molecules. Interestingly, the molecules involved in signal transduction events in both B and T lymphocytes, as well as in other non-lymphoid cells, both  
15 hematopoietic and non-hematopoietic, often bear an internal domain with homology to a domain in the *src* p60 tyrosine kinase found in platelets and neuronal tissues. This domain, termed an SH2 domain, has been found to bind to phosphorylated tyrosine residues, thus mediating the interaction of molecules involved in the signal transduction cascade in a variety of cells.

The Signaling Lymphocyte Activation Molecule (SLAM) (CDw150), a 70kD  
20 glycosylated type I transmembrane protein present on the surface of B and T cells, is a high-affinity self-ligand. Since triggering of SLAM co-activates T or B lymphocyte responses, it is considered to play a major role in bi-directional T $\leftrightarrow$ B cell stimulation (Cocks *et al.*, Nature 376: 260-263, 1995; Aversa *et al.*, Immunol. Cell Biol. 75: 202-205, 1997; Aversa *et al.*, J. Immunol. 158: 4036-4044, 1997; Carballido *et al.*, J.  
25 Immunol. 159: 4316-4321, 1997; Isomaki *et al.*, J. Immunol. 159: 2986-2993, 1997; Punnonen *et al.*, J. Exp. Med. 185: 993-1004, 1997; Ferrante *et al.*, J. Immunol. 160: 1514-1521, 1998). Moreover, triggering by anti-SLAM antibodies induces the interferon- $\gamma$  gene and redirects Th2 responses of antigen-specific T cell clones to a Th1 or Th0 phenotype (Aversa *et al.*, Immunol. Cell Biol. 75: 202-205, 1997; Aversa

-2-

*et al.*, J. Immunol. 158: 4036-4044, 1997; Carballido *et al.*, *supra*). Furthermore, since anti-SLAM antibodies alone fail to enhance B cell proliferation, and produce, if anything, an inhibitory effect (Punnonen *et al.*, *supra*), SLAM induced signal transduction events in T lymphocytes would appear to be different from those in B  
5 cells.

Given the important role of SH2 domain-containing protein mediated signal transduction pathways, in particular the pathways involved in the immune system, novel methods and reagents for modulating such pathways are useful in treating a myriad of human conditions involving inappropriate levels of such SH2 domain-  
10 containing protein signal transduction.

#### Summary of the Invention

Described herein is a novel T cell specific-protein, SAP (SLAM-Associated Protein), that contains an SH2-domain and a short tail. Additionally disclosed is the identification of a novel family of signal transduction regulators, namely the SAP  
15 family. SAP family member proteins block the recruitment of SH2-domain containing signal transduction molecules to a docking site in the SLAM cytoplasmic domain.

We mapped the SAP protein-encoding gene to the locus for X-linked proliferative disease (XLP), and mutations in the SAP gene were found in three XLP  
20 patients. Our findings led us to the conclusion that the absence of SAP in XLP patients affects T / B cell interactions induced by SLAM, which leads to an inability to control B cell proliferation induced by EBV infections. Our results demonstrate that the SAP protein plays a key role in immunoregulation, and that SAP family member proteins play an important role in SH2 domain-containing protein mediated  
25 signal transduction.

According, in a first aspect, the invention provides a substantially pure nucleic acid encoding a SAP polypeptide. In one embodiment, the SAP polypeptide is a fragment of the full length naturally-occurring SAP polypeptide. In another

-3-

embodiment, the nucleic acid is operably linked to a second nucleic acid. The second nucleic acid sequence may be a coding sequence and the nucleic acid operably linked to said second nucleic acid produces a fusion protein that includes the SAP polypeptide. The second nucleic acid sequence may also be a gene promoter. In  
5 another embodiment, nucleic acid comprises a mutation that results in an amino acid alteration in the SAP polypeptide. In preferred embodiments of this aspect of the invention, the nucleic acid includes a nucleic acid sequence that is substantially identical to SEQ ID NO: 3 or SEQ ID NO: 5. In other embodiments, the SAP polypeptide binds a phosphorylated tyrosine residue, or binds a non-phosphorylated  
10 tyrosine residue. In yet another embodiment, the nucleic acid includes a nucleic acid sequence encoding a naturally-occurring SH2 domain.

In another embodiment of the first aspect of the invention, the nucleic acid encodes a SAP polypeptide that has SAP biological activity. In other embodiments, the SAP polypeptide modulates SH2 domain-containing protein-mediated signal  
15 transduction, such as antigen-specific T cell activation, which may be mediated by Th1 cells.

In a second aspect, the invention provides a substantially pure SAP polypeptide. In one embodiment, the polypeptide is a fragment of the full length naturally-occurring SAP polypeptide. In another embodiment, the polypeptide is part  
20 of a fusion protein. In yet another embodiment, the polypeptide includes an amino acid sequence that has a mutation as compared to the naturally-occurring amino acid sequence of the polypeptide. In preferred embodiments of this aspect of the invention, the SAP polypeptide includes an amino acid sequence that is substantially identical to SEQ ID NO: 4 or SEQ ID NO: 6. In other embodiments, the SAP  
25 polypeptide binds a phosphorylated tyrosine residue, or binds a non-phosphorylated tyrosine residue. In yet another embodiment, the polypeptide includes a naturally-occurring SH2 domain.

In another embodiment of the second aspect of the invention, the SAP polypeptide has SAP biological activity. In other embodiments, the polypeptide

-4-

modulates SH2 domain-containing protein-mediated signal transduction, such as antigen-specific T cell activation which may be mediated by Th1 cells.

In a third aspect, the invention provides a method for detecting a disease involving aberrant SH2 domain-containing protein-mediated signal transduction in a patient that includes: (a) isolating a cell from the patient and (b) measuring the level of expression of a SAP family member polypeptide in the cell, where an alteration in the level in the patient relative to the level in a cell from a healthy control indicates the presence of a disease involving aberrant SH2 domain-containing protein-mediated signal transduction in the patient. Preferably, the disease is X-linked proliferative disease.

In a fourth aspect, the invention provides a method for detecting a disease involving aberrant SH2 domain-containing protein-mediated signal transduction in a patient that includes: (a) isolating a SAP family member polypeptide from the patient and (b) determining the amino acid sequence of the SAP family member polypeptide, where an alteration in the amino acid sequence in the patient relative to the amino acid sequence of a SAP family member polypeptide isolated from a healthy control indicates the presence of a disease involving aberrant SH2 domain-containing protein-mediated signal transduction in the patient. Preferably, the disease is X-linked proliferative disease.

In a fifth aspect, the invention provides a method for treating a disease involving aberrant SH2 domain-containing protein-mediated signal transduction in a patient that includes administering to the patient a SAP family member polypeptide or a fragment, mutant, or fusion thereof.

In various embodiments of the third, fourth, and fifth aspects of the invention, the SAP family member polypeptide includes an amino acid sequence substantially identical to SEQ ID NO: 4 or SEQ ID NO: 6. Preferably, the disease is X-linked proliferative disease. In other embodiments, the SAP family member polypeptide is EAT-2. In other embodiments, the SAP polypeptide binds a phosphorylated tyrosine residue or non-phosphorylated tyrosine residue. In another embodiment, the SH2

-5-

domain-containing protein-mediated signal transduction is antigen-specific T cell activation, which may be mediated by Th1 cells.

In a sixth aspect, the invention provides a method for identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction that includes: (a) providing a cell that includes a SAP family member-encoding gene; (b) contacting the cell with a candidate compound; and (c) monitoring expression of the SAP family member-encoding gene, where an alteration in the level of the expression of the gene in response to the candidate compound indicates the presence of a compound that modulates SH2 domain-containing protein-mediated signal transduction. In one embodiment of this aspect, the SAP family member-encoding gene encodes a SAP polypeptide. In another embodiment, the SAP family member encoding gene encodes an EAT-2 polypeptide.

In a seventh aspect, the invention provides a method for identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction that includes: (a) providing a cell including a reporter gene operably linked to a promoter from a SAP family member encoding gene; (b) contacting the cell with a candidate compound; and (c) measuring expression of the reporter gene, where an alteration in the level of the expression of the reporter gene in response to the candidate compound indicates the presence of a compound that modulates SH2 domain-containing protein-mediated signal transduction. In one embodiment of this aspect, the SAP family member-encoding gene encodes a SAP polypeptide. In another embodiment, the SAP family member encoding gene encodes an EAT-2 polypeptide.

In an eighth aspect, the invention provides a method for identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction that includes: (a) providing a cell having: (i) a reporter gene operably linked to a DNA-binding-protein recognition site; (ii) a first fusion gene capable of expressing a first fusion protein that includes a SAP family member polypeptide covalently bonded to a binding moiety that is capable of specifically binding to the

-6-

DNA-binding-protein recognition site; and (iii) a second fusion gene capable of expressing a second fusion protein that includes a SLAM polypeptide covalently bonded to a gene activating moiety; (b) exposing the cell to a candidate compound; and (c) measuring reporter gene expression in the cell, where an alteration in the level of the expression of the reporter gene in response to the candidate compound indicates the presence of a compound that modulates SH2 domain-containing protein-mediated signal transduction. In one embodiment of the eighth aspect, the cell is a yeast cell.

In a ninth aspect, the invention provides a method for identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction that includes: (a) providing a cell having: (i) a reporter gene operably linked to a DNA-binding-protein recognition site; (ii) a first fusion gene capable of expressing a first fusion protein that includes a SLAM polypeptide covalently bonded to a binding moiety that is capable of specifically binding to the DNA-binding-protein recognition site; and (iii) a second fusion gene capable of expressing a second fusion protein that includes a SAP family member polypeptide covalently bonded to a gene activating moiety; (b) exposing the cell to the compound; and (c) measuring reporter gene expression in the cell, where an alteration in the level of the expression of the reporter gene in response to the candidate compound indicates the presence of a compound that modulates SH2 domain-containing protein-mediated signal transduction. In one embodiment of this aspect of the invention, the cell is a yeast cell.

In various embodiments of the sixth, seventh, eighth, and ninth aspects of the invention, the SH2 domain-containing protein-mediated signal transduction is antigen-specific T cell activation, which may be mediated by Th1 cells, and where the alteration is an increase indicates the compound increases antigen-specific T cell activation, and the alteration is a decrease indicates the compound decreases antigen-specific T cell activation. In other preferred embodiments, the SAP family member polypeptide is substantially identical to SEQ ID NO: 4 or SEQ ID NO: 6. In other embodiments, the SAP family member polypeptide is EAT-2.

In a tenth aspect, the invention provides a method for identifying a compound



-7-

as a SAP family member mimetic that includes the steps of: (a) providing a SLAM polypeptide, or a fragment or fusion thereof bearing non-phosphorylated tyrosine residues; (b) contacting the SLAM polypeptide with a SAP family member polypeptide; (c) contacting the SLAM polypeptide and the SAP family member polypeptide with a candidate compound; and (d) measuring the level of interaction of the SLAM polypeptide with the SAP family member polypeptide, where a decrease in the level in response to the compound relative to a level not contacted with the compound indicates that the compound is a SAP family member mimetic. In one embodiment, the SLAM polypeptide is bound to a solid state substrate.

10 In an eleventh aspect, the invention provides a method for identifying a compound as a SAP family member mimetic that includes the steps of: (a) providing a SLAM polypeptide, or a fragment or fusion thereof, the SLAM polypeptide bearing non-phosphorylated tyrosine residues; and (b) contacting the SLAM polypeptide with a candidate compound, where the candidate compound binding to the SLAM polypeptide indicates that the candidate compound is a SAP family member mimetic. In one embodiment of this aspect of the invention, the SLAM polypeptide is bound to a solid state substrate.

In a twelfth aspect, the invention features a method for identifying a polypeptide that modulates SH2 domain-containing protein-mediated signal transduction that includes: (a) providing a cell having: (i) a reporter gene operably linked to a DNA-binding-protein recognition site; (ii) a first fusion gene capable of expressing a first fusion protein that includes a SAP family member polypeptide covalently bonded to a binding moiety that is capable of specifically binding to the DNA-binding-protein recognition site; and (iii) a second fusion gene capable of expressing a second fusion protein that is selected from a library that includes a polypeptide covalently bonded to a gene activating moiety that is encoded by a cDNA of the library; and (b) measuring reporter gene expression in the cell, where an increase in the reporter gene expression identifies the presence of a polypeptide that modulates SH2 domain-containing protein-mediated signal transduction. In a

-8-

preferred embodiment, the cell is a yeast cell. In another embodiment of this aspect of the invention, the SH2 domain-containing protein-mediated signal transduction is antigen-specific T cell activation, which may be mediated by Th1 cells. In yet another embodiment, the library is constructed from a cell selected from a group consisting of a T cell, a B cell, a myeloid cell, a natural killer cell, a hepatocyte, a  
5 liver cell, a thymocyte, a hematopoietic progenitor cell, a fibroblast, a muscle cell, and a neuron.

In a thirteenth aspect, the invention provides a method for modulating SH2 domain-containing protein-mediated signal transduction in a mammal that includes  
10 providing a transgene encoding a SAP family member polypeptide or fragment thereof to a cell of the mammal, where the transgene is positioned for expression in the cell. In a preferred embodiment of this aspect of the invention, the mammal is selected from a group consisting of a rodent (*e.g.*, a mouse), a primate (*e.g.*, a chimpanzee), a ruminant (*e.g.*, a cow), a pig, a horse, a sheep, and a goat.

15 In a fourteenth aspect, the invention provides a method for modulating SH2 domain-containing protein-mediated signal transduction in a mammal that includes administering to a cell of the mammal a compound which modulates SAP family member biological activity. In a preferred embodiment of this aspect of the invention, the mammal is selected from a group consisting of a rodent (*e.g.*, a mouse), a primate  
20 (*e.g.*, a chimpanzee), a ruminant (*e.g.*, a cow), a pig, a horse, a sheep, and a goat.

In a fifteenth aspect, the invention provides a method for increasing antigen-specific T cell activation in a mammal that includes providing a transgene encoding a SAP family member polypeptide, the transgene being positioned for expression in the cell. In a preferred embodiment of this aspect of the invention, the mammal is  
25 selected from a group consisting of a rodent (*e.g.*, a mouse), a primate (*e.g.*, a chimpanzee), a ruminant (*e.g.*, a cow), a pig, a horse, a sheep, and a goat. Preferably, the antigen-specific T cell activation is mediated by Th1 helper T cells.

In a sixteenth aspect, the invention provides a method for increasing antigen-specific T cell activation in a mammal that includes administering to a cell of the

-9-

mammal a compound which increases the biological activity of a SAP family member protein. In a preferred embodiment of this aspect of the invention, the mammal is selected from a group consisting of a rodent (*e.g.*, a mouse), a primate (*e.g.*, a chimpanzee), a ruminant (*e.g.*, a cow), a pig, a horse, a sheep, and a goat. Preferably, the antigen-specific T cell activation is mediated by Th1 helper T cells.

In a seventeenth aspect, the invention features a transgenic mammal having a knockout mutation in an endogenous SAP family member protein-encoding nucleic acid sequence. Preferably, the SAP family member protein is a SAP protein or is an EAT-2 protein. In a preferred embodiment of this aspect of the invention, the mammal has altered SH2 domain-containing protein signal transduction relative to a mammal lacking an alteration in wild-type SAP family member-encoding nucleic acid sequences. In another preferred embodiment of this aspect of the invention, the mammal is selected from a group consisting of a rodent (*e.g.*, a mouse), a primate (*e.g.*, a chimpanzee), a ruminant (*e.g.*, a cow), a pig, a horse, a sheep, and a goat.

In an eighteenth aspect, the invention features a transgenic mammal having an exogenous SAP family member protein-encoding nucleic acid sequence operably linked to a promoter. Preferably, the SAP family member protein is a SAP protein or is an EAT-2 protein. In a preferred embodiment of this aspect of the invention, the mammal has altered SH2 domain-containing protein signal transduction relative to a mammal lacking an exogenous SAP family member-encoding nucleic acid sequence operably linked to a promoter. In another preferred embodiment of this aspect of the invention, the mammal is selected from a group consisting of a rodent (*e.g.*, a mouse), a primate (*e.g.*, a chimpanzee), a ruminant (*e.g.*, a cow), a pig, a horse, a sheep, and a goat.

In accordance with the present invention, by purified "SAP family member," or "SAP family member polypeptide" is meant an isolated SH2 domain with SAP biological activity, particularly the ability to bind either a phosphorylated or a non-phosphorylated tyrosine residue. A SAP family member polypeptide has fewer than

-10-

40 amino acids, preferably fewer than 35 amino acids, located N-terminally or C-terminally to the SH2 domain in the native polypeptide. Preferably, the non-SH2 domain amino acids have no enzymatic activity. For example, if a fusion protein is made using, as one of the fusion partners an isolated SH2 domain excised from an SH2 domain-containing protein, the fusion protein is a SAP family member polypeptide if not more than 40 amino acids from the N-terminally and/or C-terminally located amino acids immediately adjacent to the SH2 domain in the native SH2 domain-containing protein are incorporated into the fusion protein. A preferred SAP family member polypeptide modulates signal transduction pathways involving SH2 domain-containing proteins. Exemplary SAP family member proteins are the SAP proteins and the EAT-2 proteins described herein.

By purified "SAP," "SAP protein," or "SAP polypeptide" is meant a protein or polypeptide having at least 70%, preferably at least 80%, more preferably at least 90%, and most preferably at least 95% overall identity with the amino acid sequence of human SAP (SEQ ID NO: 4) or murine SAP (SEQ ID NO: 6), as shown in Figs. 2B and 2D, respectively. Polypeptide products from splice variants of SAP gene sequences are also included in this definition. Preferably, the SAP protein is encoded by nucleic acid having a sequence which hybridizes to the nucleic acid sequence of Figs. 2A or 2C under high stringency conditions. Proteins and polypeptides localized anywhere in a cell are included in the definition. For example, a SAP polypeptide may be nucleus-localized, or membrane attached through myristoylation or palmitoylation addition.

By "SAP family member specific antibody" is meant an antibody (*i.e.*, monoclonal or polyclonal) that binds a SAP family member polypeptide, such as the human or mouse SAP proteins or EAT-2 proteins described herein. Specifically excluded from the definition is an antibody that binds an SH2 domain-containing protein, as defined below. Preferably, the SAP family member specific antibody of the invention is a rabbit polyclonal antibody-containing antisera described below.

By "SH2 domain" is meant a polypeptide domain which is defined by the

-11-

presence of "blocks" found in many proteins initially identified as having sequence identity to a domain in src tyrosine kinase that defines the second src homology (SH2) domain. Current practice in the field has determined the presence of an SH2 domain based on the close identity of "blocks" of sequences within the putative SH2 domain with "blocks" found in other SH2-domain containing proteins. Such analysis can be made using the position-based method of Henikoff and Henikoff, J. Mol. Biol. 243: 574-578, 1994; Henikoff and Henikoff, Genomics 19: 97-107, 1994. For example, shown in Fig. 4A is the SAP amino acid sequence with the positions of the five SH2 domain-hallmarking "blocks" shown in bold. In Fig. 4B, the sequences of the blocks in human SAP is shown. Each of these "blocks" has a high degree (*e.g.*, above 60%) sequence identity with a corresponding block in the SH2 domain of another SH2 domain-containing protein. Although as a entire family, the SH2 domain of one protein may show a moderate degree of identity with a second SH2 domain-containing protein; however the sequence identity across "blocks" identify an SH2 domain-containing protein.

By "SH2 domain-containing protein" or "SH2 domain-containing polypeptide" is meant a protein or polypeptide which has an SH2 domain, as well as at least 50 amino acid residues located N-terminal or C-terminal to the SH2 domain. Hence, specifically excluded from the definition of SH2 domain-containing polypeptides are SAP family member polypeptides. Preferably, the SH2 domain of an SH2-domain containing protein binds phosphorylated tyrosine residues, but does not bind non-phosphorylated tyrosine residues. The amino acids located N-terminal or C-terminal to the SH2 domain of an SH2 domain-containing protein may have functional activity (*e.g.*, another SH2 domain, an SH3 domain, a kinase domain, or a phosphatase domain).

By "SLAM" is meant the protein or polypeptide which is an isoform (*i.e.*, a gene splice variant) of SLAM, a multifunctional 70 kDa glycoprotein member of the Ig superfamily. SLAM is a high affinity self-ligand and is characterized by its rapid induction on naive T cells and B cells following activation of these cells. The four

-12-

SLAM isoforms currently known are SLAM 1 (or simply SLAM, the 70 kDa glycoprotein), SLAM2, SLAM3, and SLAM4.

By "SAP biological activity" or "SAP family member biological activity" is meant any one or more of the biological activities described herein for any of the SAP family member polypeptides described herein, including, without limitation, the ability to bind to a non-phosphorylated tyrosine residue, the ability to bind to a phosphorylated tyrosine residue, and the ability to counteract signaling activity of an SH2 domain-containing polypeptide. The non-phosphorylated tyrosine residue is preferably present in any polypeptide. Most preferably, the non-phosphorylated tyrosine residue is present in the cytoplasmic domain of the SLAM polypeptide.

By "SH2 domain-containing protein-mediated signal transduction" is meant a signal transduction event or signal transduction pathway in which an SH2 domain-containing protein plays a role. For example, the platelet derived growth factor (PDGF) receptor-mediated signal transduction pathway is an SH2 domain-containing protein-mediated signal transduction because the pathway involves at least one SH2 domain-containing protein.

By "modulating SH2 domain-containing protein-mediated signal transduction" is meant increasing (*i.e.*, enhancing) or decreasing (*i.e.*, inhibiting) the intracellular signaling involving SH2 domain-containing proteins in a given cell population relative to a control cell population not exposed to a test compound. Preferably, the increase or decrease in the given cell population exposed to a test compound is a change of at least 25%, more preferably the change is at least 50%, and most preferably the change is at least one-fold, as compared to a control cell population. SH2 domain-containing protein-mediated signal transduction may be measured by a variety of assays known in the art, including the assays described herein (*e.g.*, SH2 domain-containing protein-mediated signal transduction-mediated kinase activity, phosphatase activity, or change in phosphotyrosine proteins). It will be appreciated that the degree of modulation provided by a SAP polypeptide or a modulating compound in a given assay will vary, but that one skilled in the art can

-13-

determine the statistically significant change or a therapeutically effective change in the level of SH2 domain-containing protein-mediated signal transduction which identifies a SAP family member polypeptide, or identifies a compound which modulates SAP family member polypeptide or is a SAP family member therapeutic.

- 5 By "modulating T cell activation" or "altering T cell activation" is meant increasing (*i.e.*, enhancing) or decreasing (*i.e.*, inhibiting) the number of T cells that become stimulated by antigen via their antigen-specific receptors in a given cell population relative to a control cell population not exposed to a test compound. Preferably, the increase or decrease in the given cell population exposed to a test
- 10 compound is a change of at least 25%, more preferably the change is at least 50%, and most preferably the change is at least one-fold, as compared to a control cell population. T cell activation may be measured by a variety of assays known in the art, including the assays described herein (*e.g.*, T cell activation-mediated upregulation of T cell surface molecules CD69, CD25, and Fas Ligand). Preferably,
- 15 the cell population is selected from a group including TH1 CD4<sup>+</sup> T cells, TH2 CD4<sup>+</sup> T cells, and/or CD8<sup>+</sup> T cells. It will be appreciated that the degree of modulation provided by a SAP polypeptide or a modulating compound in a given assay will vary, but that one skilled in the art can determine the statistically significant change or a therapeutically effective change in the level of T cell activation which identifies a
- 20 SAP polypeptide, or identifies a compound which modulates SAP or is a SAP therapeutic.

- By "T cell activation" or "antigen-specific T cell activation" is meant a T cell that exhibits an activated phenotype (*e.g.*, increased expression of activation-dependent genes such as interleukin-2, CD69, CD25,  $\gamma$ -interferon, Fas Ligand) in
- 25 response to stimulation through the antigen-specific T cell receptor/CD3 complex. Antigen-specific T cell activation may be by stimulation of the T cell with a syngeneic antigen-presenting cell presenting the antigen in context with MHC class I or class I. For example, an I-A<sup>b</sup> restricted ovalbumin peptide-specific T cell may achieve antigen-specific T cell activation when stimulated with an H-2<sup>b</sup> antigen-

-14-

presenting cell expressing on its cell surface the ovalbumin peptide in context with class II MHC. Antigen-specific T cell activation may also be achieved by incubating the T cell with an antibody toward CD3 plus an antibody specific toward a costimulatory molecule (e.g., CD28). Antigen-specific T cell activation may be mediated by helper T cells of the Th1, Th2, or Th0 phenotype.

By "antigen" is meant a protein or polypeptide capable of eliciting an immune response. The antigen may be derived from any source.

By "antigen-presenting cell" is meant a cell that expresses on its cell surface MHC proteins. A preferable antigen presenting cell expresses MHC class II proteins on its cell surface, and a most preferable antigen presenting cell expresses both MHC class I and MHC class II proteins on its cell surface.

By "naturally-occurring," as used herein in reference to nucleic acids or polypeptides, is meant sequences found to occur in nature. Included in the definition are naturally-occurring mutations, homologues, isoforms, truncations, splice variants, and other naturally-occurring variants of the nucleic acid or polypeptide.

By "high stringency conditions" is meant conditions that are commonly understood in the art as highly stringent. Exemplary high stringency conditions include hybridizing conditions that employ low ionic strength and high temperature for washing. One, high stringency condition may include hybridization at about 40°C in about 2XSSC and 1%SDS, followed by a first wash at about 65°C in about 2XSSC and 1%SDS, and a second wash at about 65°C in about 1XSSC. Another preferred high stringency condition includes hybridizing at 2X SSC at 40°C with a probe length of at least 40 nucleotides. For other definitions of high stringency conditions, see Ausubel, F. *et al.*, Chapter 6, Current Protocols in Molecular Biology, John Wiley & Sons, New York, 1994 hereby incorporated by reference.

By "protein" or "polypeptide" is meant any chain of more than two amino acids, regardless of post-translational modification such as glycosylation or phosphorylation.

By "pharmaceutically acceptable carrier" means a carrier which is



-15-

physiologically acceptable to the treated mammal while retaining the therapeutic properties of the compound with which it is administered. One exemplary pharmaceutically acceptable carrier is physiological saline. Other physiologically acceptable carriers and their formulations are known to one skilled in the art and  
5 described, for example, in Remington's Pharmaceutical Sciences, (18<sup>th</sup> edition), ed. A. Gennaro, 1990, Mack Publishing Company, Easton, PA.

By "substantially identical" is meant a polypeptide or nucleic acid exhibiting at least 50%, preferably at least 70%, more preferably at least 85%, still more preferably at least 90%, and most preferably at least 95% identity to a reference amino  
10 acid or nucleic acid sequence. For polypeptides, the length of comparison sequences will generally be at least 16 contiguous amino acids, preferably at least 20 contiguous amino acids, more preferably at least 25 contiguous amino acids, and most preferably at least 35 contiguous amino acids. For nucleic acids, the length of comparison sequences will generally be at least 50 contiguous nucleotides, preferably at least 60  
15 contiguous nucleotides, more preferably at least 75 contiguous nucleotides, and most preferably at least 110 contiguous nucleotides.

Sequence identity is typically measured using sequence analysis software with the default parameters specified therein (e.g., Sequence Analysis Software Package of the Genetics Computer Group, University of Wisconsin Biotechnology Center,  
20 Madison, WI 53705). This software program matches similar sequences by assigning degrees of homology to various substitutions, deletions, and other modifications. Preferably, a polypeptide substantially identical to a reference polypeptide differs only by conservative amino acid substitutions. Conservative substitutions typically include substitutions within the following groups: glycine, alanine, valine, isoleucine, leucine;  
25 aspartic acid, glutamic acid, asparagine, glutamine; serine, threonine; lysine, arginine; and phenylalanine, tyrosine.

By "substantially pure polypeptide" is meant a polypeptide that has been separated from the components that naturally accompany it. Typically, the polypeptide is substantially pure when it is at least 60%, by weight, free from the

-16-

proteins and naturally-occurring organic molecules with which it is naturally associated. Preferably, the polypeptide is a SAP polypeptide that is at least 75%, more preferably at least 90%, and most preferably at least 99%, by weight, pure. A substantially pure SAP polypeptide may be obtained, for example, by extraction from  
5 a natural source (*e.g.*, T lymphocytes); by expression of a recombinant nucleic acid encoding a SAP polypeptide in a cellular system different from the cell from which it naturally originates (*e.g.*, SAP polypeptide expressed in bacteria); or by chemically synthesizing the protein. Purity can be measured by any appropriate method, *e.g.*, by column chromatography, polyacrylamide gel electrophoresis, or HPLC analysis.

10 By "substantially pure nucleic acid" is meant nucleic acid that is free of the genes which, in the naturally-occurring genome of the organism from which the nucleic acid of the invention is derived, flank the gene. The term therefore includes, for example, a recombinant nucleic acid which is incorporated into a vector; into an autonomously replicating plasmid or virus; or into the genomic nucleic acid of a  
15 prokaryote or eukaryote; or which exists as a separate molecule (*e.g.*, a cDNA or a genomic or cDNA fragment produced by PCR or restriction endonuclease digestion) independent of other sequences. The term also includes a recombinant DNA which is part of a hybrid gene encoding additional polypeptide sequence.

By "transgene" is meant any piece of DNA which is inserted by artifice into a  
20 cell, and becomes part of the genome of the organism which develops from that cell. Such a transgene may include a gene which is partly or entirely exogenous (*i.e.*, foreign) to the transgenic organism, or may represent a gene homologous to an endogenous gene of the organism.

By "transgenic" is meant any cell which includes a DNA sequence which is  
25 inserted by artifice into that cell and becomes part of the genome of the organism which develops from that cell. As used herein, the transgenic organisms are generally transgenic mammals (*e.g.*, rodents such as rats or mice) and the DNA (transgene) is inserted by artifice into the nuclear genome.

By "knockout mutation" is meant an alteration in the nucleic acid sequence

-17-

that reduces the biological activity of the polypeptide normally encoded therefrom by at least 80% relative to the unmutated gene. The mutation may, without limitation, be an insertion, deletion, frameshift mutation, or a missense mutation. Preferably, the mutation is an insertion or deletion, or is a frameshift mutation that creates a stop  
5 codon.

By "transformation" or "transfection" is meant any method for introducing foreign molecules into a cell. Methods for transformation and transfection include, without limitation, lipofection, calcium phosphate precipitation, retroviral delivery, electroporation, and biolistic transformation.

10 By "positioned for expression" is meant that the desired sequence (*e.g.*, a cDNA) is operably linked to one or more regulatory sequences (*e.g.*, a promoter) which directs transcription and translation of the desired sequence (*i.e.*, facilitates the production of, *e.g.*, a SAP polypeptide, a recombinant protein, or a sense or antisense RNA molecule).

15 By "operably linked" is meant that a two nucleic acids (*e.g.*, a coding sequence and a gene promoter) are connected such that the sequences have an effect upon each other, where the two sequences are not directly adjacent to each other in the naturally-occurring genome of an organism. For example, a gene promoter from a viral gene operably linked to a coding sequence from a mammalian gene will permit  
20 expression of the mammalian protein product encoded by the coding sequence when the appropriate molecules (*e.g.*, transcriptional activator proteins) are bound to the gene promoter sequence. Likewise, a fusion protein can be created by operably linking two coding sequences such that a polypeptide encoded by the first coding sequence is covalently bonded (via a peptide bond) to a polypeptide encoded by the  
25 second coding sequence.

By "promoter" is meant a minimal sequence sufficient to direct transcription. Also included in the invention are those promoter elements which are cell type-specific, tissue-specific, or inducible by external signals or agents; such elements may naturally occur in the 5' or 3' or intron sequence regions of the native gene, or may

-18-

naturally occur in the long terminal repeat (LTR) of a viral genome.

By "reporter gene" is meant any gene which encodes a product whose expression is detectable. A reporter gene product may have one of the following attributes, without restriction: fluorescence (*e.g.*, green fluorescent protein), enzymatic activity (*e.g.*, luciferase or chloramphenicol acetyl transferase), toxicity (*e.g.*, ricin), or an ability to be specifically bound by a second molecule (*e.g.*, biotin or a detectably-labeled antibody).

By "detectably-labeled" is meant any means for marking and identifying a molecule (*e.g.*, an oligonucleotide probe, a cDNA molecule, or an antibody), such that another molecule to which the detectably labeled molecule associates can be detected. Methods for detectably-labeling molecules are well known in the art and include, without limitation, radioactive labeling (*e.g.*, with  $^{32}\text{P}$  or  $^{35}\text{S}$ ) and nonradioactive labeling (*e.g.*, labeling by biotinylation or chemiluminescence with, for example, fluorescein).

By "antisense," as used herein in reference to nucleic acids, is meant a nucleic acid sequence that is complementary to the coding strand of a gene (*e.g.*, a murine or human SAP gene).

By "Th1 helper T cell" is meant a helper T cell (*i.e.*, a  $\text{CD4}^+$  T cell) which, when stimulated, for example, during an immune response, serves to activate cytotoxic T cells. Th1 cells produce a number of characteristic cytokines including interleukin-2, interleukin-12, and interferon- $\gamma$ .

By "Th2 helper T cell" is meant a helper T cell (*i.e.*, a  $\text{CD4}^+$  T cell) which, when stimulated, for example, during an immune response, serves to activate antibody-producing B cells. Th2 cells produce a number of characteristic cytokines including interleukin-4.

Other features and advantages of the invention will be apparent from the following description of the preferred embodiments thereof.

Brief Description of the Drawings

Fig. 1 shows the nucleic acid sequence (SEQ ID NO: 1; above) and amino acid sequence (SEQ ID NO: 2; below) of the cytoplasmic domain of SLAM.

Figs. 2A-2D shows the DNA and amino acid sequences of human and murine SAP. Fig. 2A shows the DNA sequence of human SAP (SEQ ID NO: 3). Fig. 2B shows the amino acid sequence of human SAP (SEQ ID NO: 4). Fig. 2C shows the cDNA sequence of murine SAP (SEQ ID NO: 5). Fig. 2D shows the amino acid sequence of murine SAP (SEQ ID NO: 6).

Fig. 3 shows the DNA and protein sequences of the various SAP mutants described herein (SEQ ID Nos: 7-13). Note that in the exon 2 genomic mutant, G is a point mutation identified in A1 patient. In normal individuals, it is a C. Upper case letters in the exon 2 genomic mutant represent exon2 sequence, while lower case letters represent intronic sequence.

Figs. 4A and 4B demonstrate that human SAP contains an SH2 domain. Fig. 4A shows the blocks (in bold-face type) identifying the SH2 domain in the amino acid sequence of human SAP. Fig. 4B shows the five individual SH2 domain-identifying blocks in human SAP.

Fig. 5 shows a comparison of the deduced amino-acid sequences of human SAP and murine SAP with other SH2 domain-containing proteins. Consensus  $\alpha$ -helices and  $\beta$ -sheets are indicated in brackets. Exon/intron boundaries, as derived from the murine SAP gene, are demarcated by  $\downarrow$  arrows. Four truncation mutants were generated in the tail of human SAP (del 1 to 4), and their locations in the amino acid sequence are shown with  $\uparrow$  arrows.

Figs. 6A-6D are a series of Western blotting (WB) analyses showing the

-20-

characterization of the human and murine SAP proteins and their interactions with SLAM. In Fig. 6A, human T cell tumor cells (Jurkat) and human peripheral blood lymphocytes were lysed and postnuclear lysates (1mg/ml) were immunoprecipitated with a rabbit antiserum directed at human SAP or with a control serum (3  $\mu$ l).

- 5 Samples were resolved on SDS-PAGE and immunoblotted with anti-human-SAP (1/1000 dilution). In Fig. 6B, postnuclear lysates (1mg/ml) from C57Bl/6 thymocytes were immunoprecipitated with an anti-murine SAP antibody (3  $\mu$ l), resolved on SDS-PAGE and immunoblotted with anti-mouse-SAP (1/1000 dilution). In Fig. 6C, SLAM or vector transfected EL-4 cells were cell surface biotinylated and lysed in
- 10 detergent. Postnuclear lysates (1 mg/ml) were incubated with 5  $\mu$ g of GST or GST-SAP for 1 hour in the presence of glutathione beads or 1  $\mu$ g of anti-SLAM antibodies (2E7) for 3 hours in the presence of protein G beads. Bead associated proteins were extracted, SDS-PAGE resolved, and immunoblotted with streptavidin-HRP. In Fig. 6D, human peripheral blood lymphocytes (PBL) were activated with PHA for 5 days,
- 15 and then lysed ( $1 \times 10^8$ /ml) and immunoprecipitated with an anti-SLAM monoclonal antibody (2E7) or with a control antibody. Immunoprecipitates were resolved on SDS-PAGE and immunoblotted with a anti-SLAM serum (upper panel) or with a rabbit anti-SAP antibody (lower panel).

- Figs. 7A-7C are Northern blotting analyses showing SAP expression using a
- 20  $^{32}$ P radiolabelled human SAP probe. Fig. 7A shows a Northern blotting analysis of poly A+ mRNA isolated from various human tissues (Clontech). A human  $\beta$ -actin probe was used as loading control. Fig. 7B shows a Northern blotting analysis of total RNA from different human and mouse cell lines (20  $\mu$ g/lane). A human 18S rRNA probe was used as a loading control. Fig. 7C shows a Northern blotting analysis of
- 25 human T cell subsets and two EBV+ human B cell lines. A human  $\beta$ -actin probe was used as loading control.

Figs. 8A and 8B show SAP mRNA analyses of XLP patients. Fig. 8A is a

-21-

photograph of a 10% polyacrylamide gel in which are resolved cDNAs isolated using RT PCR (reverse transcriptase-PCR) from PBMC (peripheral blood mononuclear cells) from XLP patients and healthy donors. Samples A1, B1 and B2 are from XLP patients; sample B3 is from a healthy brother of B1 and B2; and samples CT-1 and

5 CT-2 are from healthy controls. Fig. 8B is a Northern blotting analysis of total RNA from human T cell tumor cells (Jurkat), from a subset of CD45RA<sup>High</sup> cells from a healthy donor, and from PBMC from patient B2 (20 µg/lane). Specific RNA were detected using a <sup>32</sup>P radiolabelled human SAP probe; β-actin probe was used as a loading control.

10 Figs. 9A and 9B show the nucleic acid and amino acid sequences of SAP isolated from a patient with X-linked proliferative disease (XLP). DNA products were cloned in the pCR2.1 vector, and the nucleotide sequence determined on two cDNA clones in both directions using an ABI prism 377 DNA sequencer. Fig. 9A shows a comparison of the nucleic acid sequence of two cDNA clones isolated by RT-

15 PCR from an XLP patient (A1-1 and A1-2) with those of human SAP (hSAP and hSAPΔ55). Fig. 9B shows a comparison of the predicted amino acid sequences of SAP protein isolated from an XLP patient and of SAP protein isolated from a normal human. SAP.

Figs. 10A-10C show genomic analysis of patients with X-linked proliferative

20 disease. Fig. 10A depicts the location of the point mutation near exon 2 of XLP patient A1, as well as a mechanism to explain the generation of the variant form of SAP, hSAPΔ55, found in all healthy individuals. Fig. 10B is a photograph of a 10% polyacrylamide gel in which are resolved Mnl I-digested PCR amplified products of Exon 2 sequence from genomic DNA from patient A1, three healthy individuals (CT-

25 1, CT-2, and B3), and two human cell lines (Raji and Jurkat). In addition, PCR products from 78 healthy women and 30 healthy men were analyzed. Fig. 10C is a photograph of a 2% agarose gel in which are resolved PCR amplified products of

-22-

hSAP Exon 1, Exon 2, Exon 3, Exon 4, and BRCA1 Exon 2 from genomic DNA from patient A1, B1, B2, B3, and from the cell line Raji.

Figs. 11A and 11B show the CD8SLAM fusion proteins. Fig. 11A shows schematic diagrams of SLAM3, SLAM4, CD8-SLAM3 fusion protein, CD8-SLAM4 del 1 fusion protein, and CD8-SLAM4 fusion protein. Tyrosine residues in the cytoplasmic domains of these proteins are as indicated. Fig. 11B shows the amino acid sequences of the cytoplasmic domains of SLAM3 and SLAM4, as well as the amino acid sequence of a truncated cytoplasmic domain of the SLAM4 del 1 protein.

Figs. 12A-12D are a series of Western blotting (WB) analyses showing that the SAP SH2-domain binds to a specific sequence in the cytoplasmic domain of SLAM. In Fig. 12A, COS-7 cells were transfected with the indicated constructs, biotinylated, and lysed. Postnuclear lysates were immunoprecipitated with anti-CD8 (OKT8) or anti-SLAM antibodies, and immunoprecipitates were immunoblotted with anti-human-SAP rabbit sera (upper panel) or streptavidin (lower panel). In Fig. 12B, postnuclear mouse thymocytes lysates (1mg/ml) were incubated for 1 hour with the indicated peptides coupled to beads (7  $\mu$ M) or control beads in the absence or presence of free peptides (280  $\mu$ M). Bead-associated proteins were extracted and immunoblotted with anti-human SAP rabbit serum (1/1000 dilution). In Fig. 12C, postnuclear mouse thymocytes lysates (1 mg/mL) were incubated for 1 hour with SLAM Y1 peptide coupled to beads (7 mM) in the absence or presence of free peptides (70  $\mu$ M). Bead associated proteins were extracted and immunoblotted with an anti-murine-SAP rabbit serum (1/1000 dilution). In Fig. 12D, COS-7 cells were co-transfected with CD8-SLAM3 construct and one of the following SAP deletion mutants cloned into the pCMV2-FLAG construct: Del 1, Del 2, Del 3 or Del 4 (see Fig. 5). Cell surface proteins were biotinylated, and cells were lysed 48 hours after transfection. Postnuclear lysates were immunoprecipitated with 1  $\mu$ g of anti-CD8 antibody (OKT8) and immunoprecipitates were immunoblotted with anti-FLAG



-23-

antibody (KODAK, 1/1000 dilution) (upper panel) or streptavidin (lower panel).

Figs. 13A-13D are a series of Western blotting (WB) analyses showing that SAP blocks recruitment of the tyrosine phosphatase SHP-2 to the phosphorylated cytoplasmic domain of SLAM. COS-7 cells were transfected as indicated, cell surface biotinylated and lysed. In Fig. 13A, postnuclear lysates were immunoprecipitated with anti-CD8 antibodies to isolate the CD8-SLAM3 chimera, and immunoprecipitates were immunoblotted with either streptavidin (upper panel) or anti-FLAG antibody (lower panel). In Fig. 13B, postnuclear lysates were incubated for 1 hour with the SLAM Y1 peptide coupled to beads (7 $\mu$ M). Bead-associated proteins were extracted and immunoblotted with anti-FLAG antibody (1/1000 dilution). In Figs. 13C and 13D, COS-7 cells were co-transfected with a combination of SLAM, SAP, and *c-fyn* constructs. Cell surface expressed proteins were biotinylated 48 hours after transfection, and cells were lysed. Postnuclear lysates were immunoprecipitated with anti-SLAM antibodies and immunoprecipitates were immunoblotted with anti-Phosphotyrosine-HRP (Zymed Laboratories, San Francisco, CA), streptavidin-HRP (Zymed), rabbit anti-SHP2 (Santa Cruz Biotech., Santa Cruz, CA) or anti-human SAP rabbit sera. Note that in these figures, SLAM is seen as a broad band because of its extensive glycosylation (Cocks *et al.*, *supra*).

Fig. 14 is a bar graph demonstrating that SAP has a positive affect in the SLAM co-stimulatory pathway. Jurkat human T cells were transfected with an IL-2-promoter luciferase reporter construct plus SLAM, SAP, SLAM+SAP constructs, or vector only (pCDNA3). Luciferase activity was measured after stimulation with antibodies as indicated (hatched bar, no antibody; black bar, anti-SLAM, white bar; anti-CD3 plus anti-SLAM, gray bar).

Fig. 15 shows the DNA sequence of the 5' region/exon1/intron1 of the murine SAP gene (SEQ ID NO: 14).

-24-

Fig. 16 shows the DNA sequence of exon2/intron2 of the murine SAP gene (SEQ ID NO: 15).

Fig. 17 shows the DNA sequence of intron2/exon3/intron3/exon4/intron4 of the murine SAP gene (SEQ ID NO: 16).

Fig. 18 is a schematic diagram showing the map locations of the four murine SAP exons on the murine X chromosome.

Fig. 19 is a schematic diagram showing putative transcription factor binding sites on the human and murine SAP gene sequences.

Fig. 20 is a Northern blotting analysis using as a probe radiolabelled murine SAP cDNA showing the level of murine SAP mRNA species (the 0.9 kB band) in murine Th1 helper T cells (as identified by the presence of IFN- $\gamma$  mRNA) or Th2 helper T cells (as identified by the presence of IL-4 mRNA) following 6 hours of stimulation with anti-CD3 antibody plus human IL-2. Equivalent loading of the lanes is shown in the equivalent amounts of the 2.2 kB  $\beta$ -actin mRNA per lane.

Fig. 21 is a Western blotting analysis of anti-CD8 immunoprecipitations from externally biotinylated COS cells that had been transiently transfected with CD8SLAM3 + empty vector; CD8SLAM3 + Flag-tagged EAT-2, and CD8SLAM3 + Flag-tagged SAP. The upper panel shows the results obtained by Western blotting with anti-FLAG antibody; the middle panel shows the results obtained by blotting with anti-SAP antibody; and the lower panel shows blotting with streptavidin.

#### Detailed Description

We have identified the T cell protein, SAP, as the gene that is altered in patients with X-linked proliferative disease (XLP). Not only has SLAM been

-25-

discovered to be the cell surface protein to which the XLP gene product (*i.e.*, SAP) binds, but a novel T cell signal transduction pathway initiated by the co-receptor molecule SLAM has also been revealed. Furthermore, we have identified the SAP family of proteins involved in signal transduction regulation, which includes not only the novel SAP protein described herein, but also the EAT-2 protein (Thompson *et al.*, Oncogene 13: 2649-2658, 1996).

Binding of a SAP family member protein to the cytoplasmic domain of SLAM blocks the recruitment of SHP-2. Therefore two modes of SLAM signaling are likely to exist: one in which the inhibitor SAP family member acts as a negative regulator and another in which the SHP-2 dependent signal transduction pathway becomes operational. A switch between these two signalling scenarios could occur upon T cell activation, because the level of expression of the SAP protein-encoding gene was observed to diminish rapidly after triggering of the T cell receptor. It is also possible that upon activation of T cells, SAP is released from SLAM and subsequently binds to other protein(s). The concept that SAP family members act as an inhibitory molecules of SH2-domain interactions can, in general, be extended to molecules other than SLAM. The studies described herein support the model that SAP family members control signal transduction pathways that are initiated by interactions between SLAM molecules on the interface between T and B cells.

The failure of the immune system to eliminate EBV-infected B cells in XLP appears not to be caused by a B-cell-specific defect, but rather by defective EBV specific helper T cell and cytotoxic T cell responses (Lanyi *et al.*, Genomics 39: 55-56, 1997, Article Number GE964466; Purtilo, D.T., Immunology Today 4: 291-295, 1983; Seemayer *et al.*, Pediatr. Res. 38: 471-478, 1995; Sutkowski *et al.*, J. Exp. Med. 184: 971-800, 1996). As the SAP protein (*i.e.*, one of the two members of the SAP family) is found predominantly in thymus derived lymphocytes, mutations in SAP protein will most likely affect SLAM induced signal transduction events in T lymphocytes. Many XLP patients display impaired interferon- $\gamma$  production by helper T cells, suggesting a Th2 like phenotype.

-26-

Because engagement of SLAM during antigen-specific T-cell stimulation has been shown to induce IFN- $\gamma$  production and to redirect the Th2 phenotype to a Th1/Th0 phenotype (Carballido *et al.*, J. Immunol. 159: 4316-4321, 1997), insufficient responses of the T helper subset in XLP patients, could result from an impaired function of the SLAM / SAP pathway. Alternatively, since reduced EBV specific cytotoxic T cell responses have been reported in XLP patients, it is possible that EBV infected B cells cannot be "licensed" to activate T-killer cells by helper T cells that lack SAP (Ridge *et al.*, Nature 393: 474-478, 1998; Schoenberger *et al.*, Nature 393: 480-483, 1998; Bennett *et al.*, Nature 393: 478-480, 1998). In the latter model, the SLAM/SLAM binding in T/B lymphocyte interactions might serve the same purpose as CD40/CD40L interactions in activation of dendritic cells by helper T cells. Because of the results presented herein, and, in particular, because of the correlation between the XLP phenotype and the lack of binding of the respective SAP family member protein mutants to its docking site on SLAM, we conclude that a dysfunction in SLAM/SAP family member induced signal transduction pathways is likely to be the underlying cause for the inability to generate an effective T cell response capable of sustained elimination of EBV infected B cells in XLP patients.

Uncontrolled B cell proliferation in XLP patients frequently leads to non-Hodgkin's lymphomas induced by EBV. This type of non-Hodgkin lymphoma has been detected with increasing frequencies in AIDS patients, bone marrow transplant patients, and individuals who undergo T cell suppressive treatment for other reasons. Some of these disease states might be caused by abnormal regulation of a SAP family member protein-encoding gene and/or the T cell subsets that express SAP protein.

#### I. SAP Family Member Proteins' Biological Activity

SAP protein was identified by virtue of its ability to bind to the cytoplasmic tail of SLAM4 in a yeast two-hybrid screen. Interestingly, although tyrosine residues which are normally phosphorylated in mammalian cells are not phosphorylated in yeast cells, an SH2 domain is present in SAP protein. However, as the SH2 domain

-27-

in SAP protein was also found to be able to bind phosphorylated tyrosine residues, SAP binds to both non-phosphorylated and phosphorylated tyrosine residues.

Tyrosine phosphorylation of SLAM gives rise to its being bound by the protein tyrosine phosphatase, SHP2 (also known as SHPTP-2 or syp-2). The SLAM-SHP2 complex can act as a negative regulator of signal transduction cascades (Marengere *et al.*, Science 272: 1170-1173, 1996). Binding of non-tyrosine phosphorylated SLAM by SAP protein prevents SHP2 from binding and, hence, enhances SLAM's contribution to antigen-specific T cell activation.

Using SAP protein as the prototype, we also identified another member of the SAP family, namely EAT-2 (Thompson *et al.*, Oncogene 13: 2649-2658, 1996) which also binds the cytoplasmic tail of SLAM and, thus, modulated SH2 domain-containing protein-mediated signal transduction.

While antigen-specific T cell activation is enhanced by the upregulation of the expression of a SAP family member protein, given the unique ability of a SAP family member protein to bind both phosphorylated tyrosine residues and non-phosphorylated tyrosine residues, expression of SAP family member protein in a non-T cell may either enhance or inhibit SH2 domain-containing protein-mediated signal transduction in that cell. Whether expression of a SAP family member enhances or inhibits SH2 domain-containing protein-mediated signal transduction in any particular cell depends upon which SH2 domain-containing proteins are involved in any particular signalling pathway.

Identification of the SAP family allows the characterization and exploitation of SAP family members' biological activity which is useful for modulating SH2 domain-containing protein-mediated signal transduction cellular events. For example, administration of a SAP family member protein, or polypeptide fragment thereof, may enhance antigen-specific T cell activation, as measured by antigen-specific T cell activation assays known in the art and described herein. An antigen-specific T cell activation-inhibiting amount of a SAP reagent (*e.g.*, a compound that reduces the biological function of a SAP family member, such as an anti-SAP protein neutralizing

-28-

antibody or SAP protein antisense nucleic acid) may be similarly assessed. Such assays may be carried out in a cell which either expresses endogenous SAP family member proteins, or a cell into which is introduced an ectopic SAP family member polypeptide. Preferably, the cell is a T cell that is capable of undergoing antigen-specific T cell activation.

Another approach to characterize and exploit the activity of a SAP family member (*e.g.*, SAP protein) in antigen-specific T cell activation utilizes the activation of the nuclear transcription factor NFAT in TCR-mediated signal transduction. In this system the role of a SAP family member protein in NFAT activation may be readily elucidated in various assays known to the skilled artisan. For example, one method of rapidly measuring NFAT activity is through the use of a reporter gene whose expression is directed by a NFAT binding site containing promoter (Rooney *et al.*, Mol. Cell. Biol. 15: 6299-6310 1995; Luo *et al.*, J. Exp. Med. 184: 141-147, 1996). The expression vector is preferably inserted by artifice into a cell capable of undergoing antigen-specific T cell activation or is responsive to TCR-mediated signal transduction. By detecting a change in the level of expression of the reporter gene, an NFAT-inducing or inhibiting ability of a SAP family member polypeptide or reagent may be readily assessed.

These analyses may be undertaken with human or mouse SAP or EAT-2, or other SAP family member protein (*e.g.*, an isolated SH2 domain). Likewise, SAP family members' biological activity may be assayed in any cell type that has an SH2 domain-containing protein-mediated signal transduction pathway. For example, ligation of platelet derived growth factor (PDGF) with the PDGF receptor present on a fibroblast results in activation of the protein tyrosine kinase activity of the PDGF receptor. An effect of an introduced SAP family member polypeptide on PDGF receptor-mediated signaling can be assessed by a number of different ways, including changes in fibroblast cell cycle, and changes in the association of a number of SH2 domain-containing proteins with the phosphorylated PDGF receptor, such as phosphatidylinositol 3-kinase (PI-3 kinase) and phospholipase C- $\gamma$  (PLC- $\gamma$ ).

-29-

The ability of SAP family member proteins to modulate SH2 domain-containing protein-mediated signal transduction can be defined in *in vitro* systems in which alterations SH2 domain-containing protein-mediated signal transduction can be detected. For example, once a SAP family member polypeptide is ectopically  
5 expressed in a cell, SH2 domain-containing protein-mediated signal transduction can be induced in that cell by standard methods, for example, by addition of ligand to a cell whose SH2 domain-containing protein-mediated signal transduction pathway is initiated by that the ligand receptor on the cell surface (*e.g.*, addition of PDGF to a PDGF receptor-expressing fibroblast). As a control, cells are cultured under the same  
10 conditions as those induced to undergo SH2 domain-containing protein-mediated signal transduction, but either not transfected, or transfected with a vector that lacks a SAP family member-encoding insert. The ability of each SAP family member encoding construct to enhance or inhibit SH2 domain-containing protein-mediated signal transduction upon expression can be quantified by calculating the activation  
15 profiles of the cells, *i.e.*, the ratio of the increase in the number of transfected cells to the increase in the number of control cells. These experiments can confirm the presence of SH2 domain-containing protein-mediated signal transduction modulating activity of a full length SAP family member protein protein. These assays may also be performed in combination with the application of additional compounds in order to  
20 identify compounds that modulate SH2 domain-containing protein-mediated signal transduction via expression of the SAP family member protein.

## II. Characterization, Identification, and Synthesis of SAP Family Member Proteins and Polypeptides

SAP family member polypeptides may be synthesized by introducing the  
25 polypeptide-encoding nucleic acid sequences, or fragments thereof, into various cell types or using *in vitro* extracellular systems. The function of SAP family member proteins may then be examined under different physiological conditions. For example, a SAP family member polypeptide-encoding cDNA sequence may be

-30-

manipulated to characterize the expression of a given SAP family member in a particular cellular compartment. Alternatively, cell lines may be produced which over-express the SAP family member gene product allowing purification of SAP family member polypeptides for biochemical characterization, large-scale production, antibody production, and patient therapy (*e.g.*, therapy of XLP patients).

For protein expression, eukaryotic and prokaryotic expression systems may be generated in which SAP family member encoding nucleic acid sequences are introduced into a plasmid or other vector which is then used to transform living cells. Constructs in which SAP family member encoding nucleic acids containing the entire open reading frames are inserted in the correct orientation into an expression plasmid may be used for protein expression. Alternatively, portions of the SAP family member gene sequences, including wild-type or mutant SAP family member sequences, may be inserted. Prokaryotic and eukaryotic expression systems allow various important domains of the SAP family member proteins to be recovered as fusion proteins and then used for binding, structural and functional studies, and also for the generation of appropriate antibodies. Since expression of a SAP family member protein modulates signal transduction involving SH2 domain-containing proteins, it may be desirable to put the expression of a SAP family member protein under the control of an inducible promoter.

Typical expression vectors contain promoters that direct the synthesis of large amounts of mRNA corresponding to the inserted SAP family member-encoding nucleic acid in the plasmid bearing cells. They may also include eukaryotic or prokaryotic origin of replication sequences allowing for their autonomous replication within the host organism, sequences that encode genetic traits that allow vector-containing cells to be selected for in the presence of otherwise toxic drugs, and sequences that increase the efficiency with which the synthesized mRNA is translated. Stable long-term vectors may be maintained as freely replicating entities by using regulatory elements of, for example, viruses (*e.g.*, the OriP sequences from the Epstein Barr Virus genome). Cell lines may also be produced which have integrated



-31-

the vector into the genomic DNA, and in this manner the nucleic acid product is produced on a continuous basis.

Expression of foreign sequences in bacteria such as *Escherichia coli* requires the insertion of the SAP family member encoding nucleic acid sequence into a bacterial expression vector. This plasmid vector contains several elements required for the propagation of the plasmid in bacteria, and expression of inserted DNA of the plasmid by the plasmid-carrying bacteria. Propagation of only plasmid-bearing bacteria is achieved by introducing in the plasmid selectable marker-encoding sequences that allow plasmid-bearing bacteria to grow in the presence of otherwise toxic drugs. The plasmid also bears a transcriptional promoter capable of producing large amounts of mRNA from the cloned DNA. Such promoters may or may not be inducible promoters which initiate transcription upon induction. The plasmid also preferably contains a polylinker to simplify insertion of the DNA in the correct orientation within the vector.

*In vitro* expression of SAP family member proteins, fusions, polypeptide fragments, or mutants encoded by cloned DNA is also possible using the T7 late-promoter expression system in *E. coli* (see standard protocol in, *e.g.*, Ausubel *et al.*, Current Protocols in Molecular Biology, John Wiley & Sons, New York, NY, 1994). Using this system, large amounts of mRNA corresponding to the cloned SAP family member encoding DNA can be produced, and the resulting mRNA or protein can be radioactively labeled according to standard techniques.

Once the appropriate expression vectors containing a SAP family member encoding nucleic acid, or fragment, fusion, or mutant thereof, are constructed, they may be introduced into an appropriate host cell. The precise host cell used is not critical to the invention. A wide variety of expression systems may be used to produce the recombinant SAP family member proteins. Thus, sequences encoding SAP family member proteins may be introduced into a prokaryotic host (*e.g.*, *E. coli*) or in a eukaryotic host (*e.g.*, *S. cerevisiae*, insect cells such as Sf9 cells, or mammalian cells such as COS, NIH 3T3, Daudi, Jurkat, or HeLa cells). These cells

-32-

are commercially available from, for example, the American Type Culture Collection (ATCC), Manassas, VA. The method of transformation and the choice of expression vehicle (*e.g.*, expression vector) will depend on the host system selected. For example, for expression in insect cells, the baculovirus system may be used (as commercially available from, *e.g.*, Clontech, Palo Alto, CA); for expression in mammalian cells, the vaccinia virus system may be used (as described, *e.g.*, in Ausubel *et al.*, Current Protocols in Molecular Biology, John Wiley & Sons, New York, NY, 1994). If desired, the expression system may be used in conjunction with other protein expression techniques, for example, the myc tag approach described by Evan *et al.* (Mol. Cell Biol. 5:3610-3616, 1985).

Eukaryotic expression systems permit appropriate post-translational modifications to expressed SAP family member proteins, or mutants or fragments thereof. Thus, expression in eukaryotic cells enables the study of the function of the normal complete protein, specific portions of the protein, or of naturally-occurring polymorphisms and artificially produced mutated proteins. Eukaryotic cell expression also allows the identification of regulatory elements located in the 5', 3', or intronic regions of SAP family member genes, and their roles in tissue regulation of protein expression. For example, SAP family member proteins may be produced by a transiently-transfected or stably-transfected mammalian cell line. A number of vectors suitable for stable transfection of mammalian cells are available to the public (*e.g.*, see Pouwels *et al.*, Cloning Vectors: A Laboratory Manual, 1985, Supp. 1987), as are methods for constructing such cell lines (see *e.g.*, Ausubel *et al.*, *supra*). Eukaryotic cells expressing SAP family member proteins may also be used to test the effectiveness of pharmacological agents on, for example, SAP family member-associated antigen-specific T cell activation, or as means by which to study SAP family member proteins as components of other SH2 domain-containing protein-mediated signal transduction systems.

Once the recombinant protein is expressed, it can be isolated from the expressing cells by cell lysis followed by protein purification techniques, such as

-33-

affinity chromatography. In this example, an anti-SAP family member antibody, which may be produced by the methods described herein, can be attached to a column and used to isolate the recombinant SAP family member proteins. Lysis and fractionation of SAP protein-harboring cells prior to affinity chromatography may be performed by standard methods (see *e.g.*, Ausubel *et al.*, *supra*). Once isolated, the recombinant protein can, if desired, be purified further by *e.g.*, by high performance liquid chromatography (HPLC; *e.g.*, see Fisher, Laboratory Techniques In Biochemistry And Molecular Biology, Work and Burdon, Eds., Elsevier, 1980).

The SAP family member-encoding nucleic acid sequences can be altered using procedures known in the art, such as restriction endonuclease digestion, DNA polymerase fill-in, exonuclease deletion, terminal deoxynucleotide transferase extension, ligation of synthetic or cloned DNA sequences, and site-directed sequence alteration using specific oligonucleotides together with PCR. Polypeptides of the invention, particularly short SAP family member polypeptide fragments, such as the fragment corresponding to the SAP protein SH2 domain of approximately 100 amino acids in length, can also be produced by chemical synthesis (*e.g.*, by the methods described in Solid Phase Peptide Synthesis, 2nd ed., 1984, The Pierce Chemical Co., Rockford, IL). These general techniques of polypeptide expression and purification can also be used to produce and isolate useful SAP family member polypeptide fragments or analogs, as described herein.

### III. SAP Family Member Polypeptide Fragments

Polypeptide fragments including various portions of SAP family member proteins are useful in identifying the domains important for the biological activities of SAP family member proteins. Such fragments may be generated by expression of SAP family member polypeptide fragment encoding nucleic acid fragments, generated using the nucleotide sequences provided herein, or by chemical synthesis.

SAP family member polypeptide fragments are useful, for example, in evaluating the portions of the SAP family member protein (*e.g.*, the SH2 domain)

-34-

involved in SH2 domain-containing protein-mediated signal transduction. In one example, polypeptide fragments of SAP family members are used to induce or prevent activation-dependent gene expression with or without antigen stimulation through the T cell receptor. T cell activation assays are known in the art, and include, without limitation, expression of activation-dependent genes, such as the cell surface receptors CD25, CD69, and FAS ligand, and production of cytokines, such as interleukin-2 (IL-2) and interferon- $\gamma$  (IFN- $\gamma$ ). Preferably, such SAP family member polypeptide fragments may include the SAP:SLAM or the EAT2:SLAM binding domain, or the SAP protein SH2 domain.

10 In another example, polypeptide fragments of SAP family members are used to induce or inhibit an SH2 domain-containing protein-mediated signal transduction event in a non-lymphoid cell. SAP family member polypeptide fragment activities are assessed using one or more known assays which depend upon the cell type and the pathway in which the SH2 domain-containing protein-mediated signal transduction event plays a role. For example, a SAP fragment-mediated modulation of cell proliferation induction by an SH2 domain-containing protein-mediated signal transduction event (*e.g.*, ligation of the epidermal growth factor (EGF) receptor by EGF) may be readily assayed in epidermal cell proliferation assays by modifying known cell proliferation techniques (see, *e.g.*, Ausubel *et al.*, *supra*). In T cells, SAP family member polypeptide fragments may also be used to inhibit normal activities of an endogenous full-length SAP family member in a cell exhibiting an inappropriately high level of activity of a SAP family member protein (*e.g.*, inappropriately high level of SAP protein activity in a T cell of a patient with an autoimmune disease).

#### IV. SAP Family Member Antibodies

25 In addition to the rabbit polyclonal antibodies that recognize human SAP protein and murine SAP described below, monoclonal antibodies directed toward SAP family member proteins (*e.g.*, human EAT-2) may be produced by using as antigen a SAP family member protein isolated from SAP family member-expressing

-35-

cultured prokaryotic or eukaryotic cells or a SAP family member isolated from expressing cells (e.g., EAT-2 from Ewing's sarcoma cells). Methods for generating monoclonal antibodies are well known in the art and are described, for example, in Kohler *et al.*, Nature 256: 495, 1975; Kohler *et al.*, Eur. J. Immunol. 6:511, 1976; 5 Kohler *et al.*, Eur. J. Immunol. 6: 292, 1976; Hammerling *et al.*, Monoclonal Antibodies and T Cell Hybridomas, Elsevier, New York, NY, 1981; and Coligan *et al.*, Current Protocols in Immunology John Wiley & Sons, New York, NY, 1994. Once produced, monoclonal antibodies may be tested for specific SAP family member protein recognition by standard Western blotting or immunoprecipitation 10 analyses.

Antibodies that specifically recognize SAP family member proteins may be produced using amino acid sequences that do not reside within highly conserved regions, and that appear likely to be antigenic, as analyzed by criteria such as those provided by the PeptideStructure Program (Genetics Computer Group Sequence 15 Analysis Package, Program Manual for the GCG Package, Version 7, 1991) using secondary structure according to either the Chou-Fasman method (Adv. in Enzymol. 47: 45-148, 1978) or the Garnier-Osguthorpe-Robson method (Garnier *et al.*, J. Mol. Biol. 120: 97, 1978); hydrophilicity according to either the Kyte-Doolittle method or Hopp-Woods method (Proc. Natl. Acad. Sci. USA 78: 3824-3828, 1981); surface 20 probability according to the Emini method (Emini *et al.*, J. Virol., 55: 836-839, 1985); flexibility according to the Karplus-Schulz method; and the antigenic index according to the Jameson-Wolf method (CABIOS 4(1): 181-186, 1988). Such SAP sequences preferably reside in the short cytoplasmic tail of SAP (*i.e.*, the final 26 amino acid residues of human SAP, the final 24 amino acid residues of murine SAP, or the final 25 32 amino acids of murine EAT-2). Conversely, antibodies may be produced using SAP amino acid sequences that reside within highly conserved regions. For example, amino acid sequences from the SH2 domain of SAP may be used as antigen to generate antibodies specific toward human or murine SAP.

In addition to intact monoclonal and polyclonal anti-SAP family member

-36-

antibodies, various genetically engineered antibodies, humanized antibodies, and antibody fragments, including F(ab')<sub>2</sub>, Fab', Fab, Fv, and sFv fragments, may generated that specifically recognize human or murine SAP protein, human or murine EAT-2 protein, or fragments thereof. Antibodies can be humanized by methods  
5 known in the art (e.g., by expression in transgenic animals, as described in Green *et al.*, Nature Genetics 7: 13-21, 1994). In addition, monoclonal antibodies with a desired binding specificity can be commercially humanized (Scotgene, Scotland; Oxford Molecular, Palo Alto, CA).

In addition, Ladner (U.S. Patent Nos. 4,946,778 and 4,704,692) describes  
10 methods for preparing single polypeptide chain antibodies. Ward *et al.* (Nature 341: 544-546, 1989) describe the preparation of heavy chain variable domains, which they term "single domain antibodies," which have high antigen-binding affinities. McCafferty *et al.* (Nature 348: 552-554, 1990) show that complete antibody V domains can be displayed on the surface of fd bacteriophage, that the phage bind  
15 specifically to antigen, and that rare phage (one in a million) can be isolated after affinity chromatography. Boss *et al.* (U.S. Patent 4,816,397) describe various methods for producing immunoglobulins, and immunologically functional fragments thereof, which include at least the variable domains of the heavy and light chain in a single host cell. Cabilly *et al.* (U.S. Patent 4,816,567) describe methods for preparing  
20 chimeric antibodies.

V. Identification of Compounds that Modulate Expression of SAP Family Member Proteins

Based on our experimental results, we have developed a number of screening procedures for identifying therapeutic compounds (e.g., that modulate SH2 domain-  
25 containing protein-mediated signal transduction) which can be used in human patients. In one particular example, compounds that upregulate expression of SAP family member proteins are considered useful for the treatment of patients with XLP. Likewise, compounds that downregulate SAP family member protein expression are

-37-

useful in treating patients with diseases hallmarked by an excessive amount of antigen-specific T cell activation (*e.g.*, autoimmune diseases). In general, the screening methods of the invention involve screening any number of compounds for therapeutically active agents by employing any number of *in vitro* or *in vivo* experimental systems.

The methods of the invention simplify the evaluation, identification, and development of active agents for the treatment and prevention of conditions involving an inappropriate level of SH2 domain-containing protein-mediated signal transduction, which may be excessive or insufficient, depending upon the condition. These screening methods provide a facile means for selecting natural product extracts or compounds of interest in a large population, which are further evaluated and condensed to a few active and selective materials. Constituents of this pool are then purified and evaluated to determine their SH2 domain-containing protein-mediated signal transduction inhibiting or inducing activities.

In general, novel drugs for the treatment of conditions involving an inappropriate level of SH2 domain-containing protein-mediated signal transduction are identified from large libraries of both natural product or synthetic (or semi-synthetic) extracts or chemical libraries according to methods known in the art. Those skilled in the field of drug discovery and development will understand that the precise source of test extracts or compounds is not critical to the screening procedure(s) of the invention. Accordingly, virtually any number of chemical extracts or compounds can be screened using the exemplary methods described herein. Examples of such extracts or compounds include, but are not limited to, plant-, fungal-, prokaryotic-, or animal-based extracts, fermentation broths, and synthetic compounds, as well as modification of existing compounds. Numerous methods are also available for generating random or directed synthesis (*e.g.*, semi-synthesis or total synthesis) of any number of chemical compounds, including, but not limited to, saccharide-, lipid-, peptide-, and nucleic acid-based compounds. Synthetic compound libraries are commercially available from Brandon Associates (Merrimack, NH) and Aldrich

-38-

Chemical (Milwaukee, WI). Alternatively, libraries of natural compounds in the form of bacterial, fungal, plant, and animal extracts are commercially available from a number of sources, including Biotics (Sussex, UK), Xenova (Slough, UK), Harbor Branch Oceanographics Institute (Ft. Pierce, FL), and PharmaMar, U.S.A. (Cambridge, MA). In addition, natural and synthetically produced libraries are produced, if desired, according to methods known in the art. Furthermore, any library or compound may be readily modified using standard chemical, physical, or biochemical methods.

In addition, those skilled in the art of drug discovery and development understand that methods for dereplication (*e.g.*, taxonomic dereplication, biological dereplication, and chemical dereplication, or any combination thereof) or the elimination of replicates or repeats of materials already known for their SH2 domain-containing protein-mediated signal transduction modulating activities should be employed whenever possible.

When a crude extract is found to have SH2 domain-containing protein-mediated signal transduction inhibiting or inducing activities, or both, further fractionation of the positive lead extract is necessary to isolate chemical constituents responsible for the observed effect. Thus, the goal of the extraction, fractionation, and purification process is the careful characterization and identification of a chemical entity within the crude extract having SH2 domain-containing protein-mediated signal transduction modulating activities. The same *in vivo* and *in vitro* assays described herein for the detection of activities in mixtures of compounds can be used to purify the active component and to test derivatives thereof. Methods of fractionation and purification of such heterogenous extracts are known in the art. Compounds shown to be useful agents for the treatment of pathogenicity may be chemically modified according to methods known in the art. Compounds identified as being of therapeutic value are subsequently analyzed using any standard animal model of degenerative disease or cancer known in the art.

Below we describe screening methods for identifying and evaluating the



-39-

efficacy of a compound as a SH2 domain-containing protein-mediated signal transduction modulating agent, and in particular, an antigen-specific T cell activation inhibiting or inducing agent. These methods are intended to illustrate, not limit, the scope of the claimed invention.

5            a) Screens for Compounds Affecting SAP Family Member Protein Expression

SAP family member protein encoding DNAs may be used to facilitate the identification of compounds that increase or decrease SAP family member protein expression. In one approach, candidate compounds are added, in varying concentrations, to the culture medium of cells expressing SAP family member  
10       mRNA. The SAP mRNA expression is then measured, for example, by Northern blotting analysis (Ausubel *et al.*, *supra*) using a SAP family member protein encoding DNA, cDNA, or RNA fragment as a hybridization probe. The level of SAP family member protein encoding mRNA expression in the presence of the candidate  
15       compound is compared to the level of mRNA expression of a SAP family member protein in the absence of the candidate compound, all other factors (*e.g.*, cell type and culture conditions) being equal.

The effect of candidate compounds on SAP family member protein modulation of SH2 domain-containing protein-mediated signal transduction may, instead, be measured at the level of translation by using the general approach  
20       described above with standard protein detection techniques, such as Western blotting or immunoprecipitation with an antibody specific toward a SAP family member protein (for example, the SAP protein-specific antibody described herein).

In an alternative approach to detecting compounds which regulate SAP at the level of transcription, candidate compounds may be tested for an ability to regulate a  
25       reporter gene whose expression is directed by a SAP family member-encoding gene promoter. For example, a cell which does not express a SAP family member may be transfected with a expression plasmid, such as a luciferase reporter gene operably linked to the SAP family member-encoding gene promoter. Candidate compounds may then be added, in varying concentrations, to the culture medium of the cells.

-40-

Luciferase expression levels may then be measured, for example, using the luciferase assay system kit used herein that is commercially available from Promega (Madison, WI), and rapidly assessing the level of luciferase activity on a luminometer. The level of luciferase expression in the presence of the candidate compound is compared to the level of luciferase expression in the absence of the candidate compound, all other factors (e.g., cell type and culture conditions) being equal.

Compounds that modulate the level of expression of a SAP family member protein may be purified, or substantially purified, or may be one component of a mixture of compounds such as an extract or supernatant obtained from cells, from mammalian serum, or from growth medium in which mammalian cells have been cultured (Ausubel *et al.*, *supra*). In an assay of a mixture of compounds, expression of a SAP family member protein is tested against progressively smaller subsets of the compound pool (e.g., produced by standard purification techniques such as HPLC or FPLC) until a single compound or minimal number of effective compounds is demonstrated to modulate expression of a SAP family member protein.

b) Screens for Compounds Affecting SAP Family Member Protein Biological Activity

Compounds may also be screened for their ability to affect the ability of a SAP family member protein to modulate, for example, SH2 domain-containing protein-mediated signal transduction. In this approach, the degree of SH2 domain-containing protein-mediated signal transduction in the presence of a candidate compound is compared to the degree of SH2 domain-containing protein-mediated signal transduction in its absence, under equivalent conditions. Again, the screen may begin with a pool of candidate compounds, from which one or more useful modulator compounds are isolated in a step-wise fashion. The level of an SH2 domain-containing protein mediated signal transduction pathway may be measured by any standard assay. In particular, the SH2 domain-containing protein-mediated antigen-specific T cell activation signaling pathway may be measured using T cell activation assays, many of which are described herein.

-41-

Another method for detecting compounds that modulate the SH2 domain-containing protein mediated signal transduction modulating activity of a SAP family member protein is to screen for compounds that interact physically with a given SAP family member polypeptide in a yeast two-hybrid system, as described below.

- 5           Another method for detecting protein interactions is an *in vitro* binding assay. In this method, one protein, which may be recombinantly produced and may be a fusion protein, is used like an antibody in an immunoprecipitation reaction to bind to and remove its interacting protein partner from a sample, such as a cell lysate. For example, the GST-SAP fusion protein described below may be recombinantly
- 10       produced in bacteria and used to isolate a compound capable of binding a SAP family member. Resolution of the bead-bound proteins by SDS-PAGE will detect a SAP family member interacting protein, which may then be sequenced by N-terminal peptide sequencing.

- A compound that promotes an increase in the expression or biological activity
- 15       of the SAP family member protein is considered particularly useful in the invention; such a molecule may be used, for example, as a therapeutic to increase cellular levels of a SAP family member and thereby exploit the ability of SAP family member polypeptides to modulate SH2 domain-containing protein mediated signal transduction. For example, a compound that can increase the level of a SAP family
- 20       member protein and thereby induce the SAP family member protein-mediated enhancement of antigen-specific T cell activation would be advantageous in the treatment of diseases involving insufficient antigen-specific T cell activation (*e.g.*, the XLP disease, tuberculosis, AIDS, schistosomiasis) or cancer (*e.g.*, breast cancer, prostate cancer, leukemia).

- 25       A compound that decreases the activity of a SAP family member protein (*e.g.*, by decreasing SAP family member protein gene expression or biological activity) may be used to decrease the ability of a SAP family member protein to modulate SH2 domain-containing protein mediated signal transduction. For example, a compound that decreases a SAP family member protein, thereby decreasing antigen-specific T

-42-

cell activation, would be advantageous in the treatment of diseases involving an excessive level of antigen-specific T cell activation, such as autoimmune diseases.

Preferred screens for compounds that affect a SAP family member protein's modulation of SH2 domain-containing protein-mediated signal transduction are rapid and high through-put. Using as an example the ability of SAP protein to enhance antigen-specific T cell activation, syngeneic antigen-presenting cells (APCs) expressing, for example, an ovalbumin peptide, may be cultured in a multiwell (*e.g.*, a 96 well microtiter) plate. T cells that specifically recognize ovalbumin peptide in context with the MHC expressed on the cultured syngeneic antigen presenting cells may be transfected with DNA that includes the promoter of a gene whose expression is upregulated upon antigen-specific T cell activation (*e.g.*, the promoter from the gene encoding IL-2) operably linked to a reporter gene, such as green fluorescent protein (GFP). The T cells are added to each peptide plus APC-containing well, followed by addition to each well of a compound, or combination thereof, being screened for an ability to modulate SAP protein-mediated enhancement of antigen-specific T cell activation. The compound-treated plate is then subjected to analysis on a 96 well fluorescent plate reader for the presence of GFP. A well with increased GFP expression compared to a control well not treated with any compound indicates a compound with an ability to modulate SAP protein-mediated enhancement of antigen-specific T cell activation.

Molecules that are found, by the methods described above, to effectively modulate gene expression of a SAP family member protein or biological activity of such a protein may be tested further in animal models. If they continue to function successfully in an *in vivo* setting, they may be used as therapeutics to either inhibit or enhance SH2 domain-containing protein-mediated signal transduction, as appropriate.

c) Screens for Additional Reagents which Mimic SAP Family Member Proteins

Different compounds may have the same mechanism of action without concomitant similarity in size and/or structure. Accordingly, although peptide and

-43-

non-peptide mimetics based upon the SH2 domain of a SAP family member protein are likely to induce an SAP family member-like modulation of SH2 domain-containing protein-mediated signal transduction in cells having such a signal transduction pathway, non-structural mimetic reagents may also have this capability.

- 5 Utilizing the SH2 domain-containing protein-mediated signal transduction assays described herein, such reagents may be identified.

d) Identification and Generation of SAP Family Member Protein SH2 Domain Peptide Mimetics

- One efficient method to treat patients (suffering from, for example, XLP  
10 disease) with the SH2 domain of a SAP family member is to generate mimetics which possess the non-phosphorylated tyrosine residue-binding abilities of the SAP family member proteins. Peptide mimetics of the SH2 domain of a SAP family member, or DNA encoding these peptides, may be used in concert. For example, two or more different peptides which correspond to different regions of the SAP protein SH2  
15 domain may be introduced into the same population of cells. Likewise, a peptide that corresponds to the SH2 domain of an SAP protein may be introduced with a peptide that corresponds to the SH2 domain of an EAT-2 protein.

- Synthetic peptides corresponding to the SH2 domain or a SAP family member protein may be purchased from commercially available sources (such as the peptide  
20 generating facility at the Department of Biochemistry and Molecular Pharmacology at Harvard Medical School, Boston, MA), and tested for an ability to affect SH2 domain-containing protein mediated signal transduction events, such as the ability of a SAP family member to inhibit antigen-specific T cell activation, in various signal transduction detecting assays known in the art and described herein. Such peptides  
25 may be taken up directly by cells in culture or delivered to the cells by a variety of methods, including lipid vesicles or electroporation. In addition, nucleic acid sequences encoding these peptides may be subcloned into the cloning site of an expression cloning vector and the plasmid DNA introduced to the cell of interest by various transfection methods known in the art (*e.g.*, electroporation, DEAE-dextran,

-44-

calcium phosphate). DNA encoding peptides corresponding to the SH2 domain of a SAP family member may also be incorporated into coding sequences of fusion proteins and the mimetic delivered by transfection of the fusion protein encoding expression vector or fusion protein encoding viral vectors.

- 5           Peptides, or combinations thereof, may be screened for efficacy and effective dose requirements using the various SH2 domain-containing protein-mediated signal transduction assays well known in the art. Such SH2 domain-containing protein-mediated signal transduction assays include *in vitro* kinase and *in vitro* phosphatase assays of SH2 domain-containing or phosphotyrosine residue-containing proteins  
10           with kinase and phosphatase activities, respectively. In another example, various concentrations of peptide may be introduced with the SAP family member promoter-luciferase plasmid described herein.

- DNA encoding potential peptide mimetics of the SH2 domain of a SAP family member may also be identified by hybridization of the DNA to the nucleotide  
15           sequences encoding SAP protein (SEQ ID NOs: 3 and 5) provided on Figs. 2A and 2C, or to the nucleotide sequences encoding EAT-2 protein (GenBank Accession Nos. AF020263 and AF020264). In one particular example of this approach, DNAs may be identified by an ability to hybridize to the SAP family member-encoding nucleotide sequences under high stringency conditions. High stringency conditions  
20           may include hybridization at about 40°C in about 2XSSC and 1%SDS, followed by a first wash at about 65°C in about 2XSSC and 1%SDS, and a second wash at about 65°C in about 1XSSC. Other hybridization stringency conditions, both high and low, are defined in the art (see, for example, Sambrook, Fritsch and Maniatis, Molecular Cloning: A Laboratory Manual (2nd ed.), CSH Press, 1989, or Ausubel, *et al.*,  
25           *supra*).

          Once identified, a DNA which hybridizes to the nucleotide sequences encoding a SAP family member may be used to generate a polypeptide product by standard techniques. For example, the DNA may be subcloned into the pCDNA3.1 expression plasmid (commercially available from Clontech) and the resulting plasmid

-45-

transfected into, for example, COS cells (commercially available from the ATCC), to produce recombinant polypeptide. This polypeptide product may then be screened for SAP biological activity using the various SH2 domain-containing protein-mediated signal transduction assays described herein.

- 5           The identification of minimal peptide killing sequences allows the generation of non-peptidic mimetics. Techniques for generating such non-peptidic mimetics of the SH2 domain of a SAP family member are standard chemistry techniques and well known to one skilled in the art of combinatorial chemistry. The efficacies of these non-peptide mimetics may be assayed similarly to peptide mimetics of the SAP
- 10   family member SH2 domain.

#### VI. Therapies

- Therapies may be designed to circumvent or overcome a SAP gene defect or inadequate SAP family member gene expression, and thus modulate and possibly alleviate conditions involving an inappropriate amount of SH2 domain-containing
- 15   protein-mediated signal transduction. Hence, in considering various therapies, such therapies may be targeted at any tissues demonstrated to express SAP family member protein (e.g., SAP protein in T cells). With particular focus on the immune system, therapies to inhibit human SAP family member gene expression are useful in reducing antigen-specific T cell activation in autoreactive T lymphocytes. Likewise, therapies
- 20   to enhance SAP family member gene expression are useful in enhancing antigen-specific T cell activation in, for example, XLP patients.

- SH2 domain-containing protein-mediated signal transduction-modulating SAP family member reagents may include, without limitation, full length or fragment SAP family member polypeptides, SAP family member antisense nucleic acid, or any
- 25   compound which increases a SAP family member's SH2 domain-containing protein-mediated signal transduction-modulating activity.

##### a) Protein Therapy

Treatment or prevention of inappropriate SH2 domain-containing protein-

-46-

mediated signal transduction can be accomplished by replacing mutant or surplus SAP family member protein with normal protein, by modulating the function of mutant protein, or by delivering normal SAP family member protein to the appropriate cells. It is also possible to modify the pathophysiologic pathway (e.g.,  
5 a signal transduction pathway involving SH2 domain containing proteins) in which the SAP protein participates in order to correct the physiological defect.

To replace a mutant protein with normal protein, or to add protein to cells which no longer express sufficient levels of a SAP family member, it is necessary to obtain large amounts of pure SAP family member protein from cultured cell systems  
10 which can express the protein. Delivery of the SAP family member protein to the affected cells can then be accomplished using appropriate packaging or administering systems. Alternatively, small molecule analogs may be used and administered to act as SAP family member protein agonists and in this manner produce a desired physiological effect. Methods for finding such molecules are provided herein.

15 b) Gene Therapy

Gene therapy is another potential therapeutic approach in which normal copies of the SAP family member encoding gene are introduced into T cells to successfully encode for normal and abundant protein in cells which undergo inappropriately low levels of T cell activation. Likewise, nucleic acids encoding SAP family member  
20 antisense RNA may be introduced into T cells to inhibit an inappropriately high level of T cell activation. SAP family member-encoding sequences or SAP family member antisense RNA may also be introduced into non-lymphoid cells, to modulate inappropriate levels of SH2 domain-containing protein-mediated signal transduction. The SAP family member-encoding sequence (or antisense SAP family member RNA-  
25 encoding sequence) must be delivered to those cells in a form in which it can be taken up and encode for sufficient protein to provide effective function. Preferably, such sequences are operably linked to the endogenous SAP family member gene promoter. Alternatively, in some mutants it may be possible to promote normal levels of SH2 domain-containing protein-mediated signal transduction by introducing another copy



-47-

of the homologous gene bearing a second mutation in that gene or to alter the mutation, or use another gene to block any negative effect.

Transducing retroviral vectors can be used for somatic cell gene therapy especially because of their high efficiency of infection and stable integration and expression. The targeted cells however must be able to divide and the expression levels of normal protein should be high. For example, the full length SAP family member gene, or portions thereof, can be cloned into a retroviral vector and driven from the endogenous SAP family member gene promoter or from the retroviral long terminal repeat or from a promoter specific for the target cell type of interest (e.g., the CD2 promoter in T cells). Other viral vectors which can be used include adenovirus, adeno-associated virus, vaccinia virus, bovine papilloma virus, or a herpes virus such as Epstein-Barr Virus.

Retroviral vectors, adenoviral vectors, adenovirus-associated viral vectors, or other viral vectors with the appropriate tropism for cells likely to be involved in SH2 domain-containing protein-mediated signal transduction (for example, T cells) may be used as a gene transfer delivery system for a therapeutic SAP family member gene construct. Numerous vectors useful for this purpose are generally known (Miller, Human Gene Therapy 15-14, 1990; Friedman, Science 244:1275-1281, 1989; Eglitis and Anderson, BioTechniques 6: 608-614, 1988; Tolstoshev and Anderson, Curr. Opin. Biotech. 1: 55-61, 1990; Sharp, The Lancet 337: 1277-1278, 1991; Cornetta *et al.*, Nucl. Acid Res. and Mol. Biol. 36: 311-322, 1987; Anderson, Science 226: 401-409, 1984; Moen, Blood Cells 17: 407-416, 1991; Miller *et al.*, Biotech. 7: 980-990, 1989; Le Gal La Salle *et al.*, Science 259: 988-990, 1993; and Johnson, Chest 107: 77S-83S, 1995). Retroviral vectors are particularly well developed and have been used in clinical settings (Rosenberg *et al.*, N. Engl. J. Med 323: 370, 1990; Anderson *et al.*, U.S. Patent No. 5,399,346).

Gene transfer could also be achieved using non-viral means requiring transfection *in vitro*. Transplantation of normal genes into the affected cells of a patient can also be useful therapy. In this procedure, a normal SAP family member

-48-

gene is transferred into a cultivatable cell type, either exogenous or endogenous to the patient. These cells are then injected serotologically into the targeted tissue(s). In addition, SAP family member-encoding DNA may be introduced into a T cell by lipofection (Felgner *et al.*, Proc. Natl. Acad. Sci. USA 84: 7413, 1987; Ono *et al.*,  
5 Neurosci. Lett. 117: 259, 1990; Brigham *et al.*, Am. J. Med. Sci. 298: 278, 1989; Staubinger *et al.*, Meth. Enz. 101: 512, 1983), asialorosonucoid-polylysine conjugation (Wu *et al.*, J. Biol. Chem. 263: 14621, 1988; Wu *et al.*, J. Biol. Chem. 264: 16985, 1989); or, less preferably, micro-injection under surgical conditions (Wolff *et al.*, Science 247: 1465, 1990).

10 In another approach that may be utilized with all of the above methods, a therapeutic SAP family member DNA construct is preferably applied to the site of the desired normal SH2 domain-containing protein-mediated signal transduction (for example, by injection). However, it may also be applied to tissue in the vicinity of the affected SH2 domain-containing protein-mediated signal transduction event or to a  
15 blood vessel supplying the cells with an inappropriate level of SH2 domain-containing protein-mediated signal transduction.

In the constructs described, SAP family member cDNA expression can be directed from any suitable promoter (*e.g.*, the human cytomegalovirus (CMV), simian virus 40 (SV40), or, preferably, the endogenous SAP gene promoter), and regulated  
20 by any appropriate mammalian regulatory element. For example, if desired, enhancers known to preferentially direct gene expression in neural cells, lymphocytes, or muscle cells may be used to direct expression of a SAP family member protein. The enhancers used could include, without limitation, those that are characterized as tissue- or cell-specific in their expression. Alternatively, if a SAP family member  
25 genomic clone is used as a therapeutic construct, regulation may be mediated by the cognate regulatory sequences or, if desired, by regulatory sequences derived from a heterologous source, including any of the promoters or regulatory elements described above.

Antisense based strategies may be employed to explore SAP family member

-49-

gene function and as a basis for therapeutic drug design. Antisense strategies may use a variety of approaches, including the use of antisense oligonucleotides and injection of antisense RNA. For our analysis of SAP family member gene function, a method of transfection of antisense RNA expression vectors into targeted cells is employed.

- 5 Phenotypic effects induced by antisense effects are based on changes in criteria such as protein levels, protein activity measurement, and target mRNA levels.

For example, SAP family member gene therapy may also be accomplished by direct administration of antisense SAP family member mRNA to a cell that is expected to undergo an undesirably high level of T cell activation (*e.g.*, an  
10 autoreactive T cell). The antisense SAP family member mRNA may be produced and isolated by any standard technique, but is most readily produced by *in vitro* transcription using an antisense SAP family member cDNA under the control of a high efficiency promoter (*e.g.*, the T7 promoter). Administration of antisense SAP family member nucleic acid to cells can be carried out by any of the methods for  
15 direct nucleic acid administration described above.

Another therapeutic approach within the invention involves administration of recombinant SAP polypeptide, either directly to the site of a desired inhibition of SH2 domain-containing protein-mediated signal transduction event (for example, by injection) or systemically (for example, by any conventional recombinant protein  
20 administration technique). The dosage of SAP family member depends on a number of factors, including the size and health of the individual patient, but, generally, between 0.1 mg and 100 mg inclusive are administered per day to an adult in any pharmaceutically acceptable formulation.

#### c) Preventative Therapies

- 25 In a patient diagnosed to have a SAP family member mutation or to be susceptible to SAP family member mutations or to have aberrant expression of a SAP family member (even if those mutations or expression patterns do not yet result in underexpression or decreased biological activity of a SAP family member), or a patient diagnosed with a XLP disease, or diagnosed with an infectious disease (*e.g.*,

-50-

AIDS), any of the above therapies may be administered before the occurrence of the disease phenotype. For example, the therapies may be provided to an XLP patient who does not yet show EBV-infection induced non-Hodgkin's lymphoma. In particular, compounds shown to increase SAP family member expression or SAP family member biological activity may be administered to patients diagnosed with infectious diseases or cancer by any standard dosage and route of administration (see above). Alternatively, gene therapy using a antisense SAP family member mRNA expression construct may be undertaken to reverse or prevent the T cell defect prior to the development of an autoimmune disease.

- 10           The methods of the instant invention may be used to reduce or diagnose the disorders described herein in any mammal, for example, humans, domestic pets, or livestock. Where a non-human mammal is treated or diagnosed, the SAP family member polypeptide, nucleic acid, or antibody employed is preferably specific for that species.

15   VII.   Administration of SAP Reagents and Modulators

- A SAP family member protein, gene, or modulator may be administered within a pharmaceutically-acceptable diluent, carrier, or excipient, in unit dosage form. Conventional pharmaceutical practice may be employed to provide suitable formulations or compositions to administer neutralizing SAP family member-specific antibodies or SAP family member-inhibiting compounds (*e.g.*, antisense SAP family member or a SAP family member dominant negative mutant) to patients suffering from a disease characterized by an excessive level of T cell activation (*e.g.*, an autoimmune disease). Likewise, a SAP family member protein, a cDNA encoding a SAP family member protein, or a mimetic thereof, may be administered to a patient suffering from a disease characterized by an insufficient level of T cell activation (*e.g.*, XLP or cancer). Administration may begin before the patient is symptomatic. Any appropriate route of administration may be employed, for example, administration may be parenteral, intravenous, intra-arterial, subcutaneous,

-51-

intramuscular, intracranial, intraorbital, ophthalmic, intraventricular, intracapsular, intraspinal, intracisternal, intraperitoneal, intranasal, aerosol, by suppositories, or oral administration. Therapeutic formulations may be in the form of liquid solutions or suspensions; for oral administration, formulations may be in the form of tablets or capsules; and for intranasal formulations, in the form of powders, nasal drops, or aerosols.

Methods well known in the art for making formulations are found, for example, in Remington's Pharmaceutical Sciences, (18<sup>th</sup> edition), ed. A. Gennaro, 1990, Mack Publishing Company, Easton, PA. Formulations for parenteral administration may, for example, contain excipients, sterile water, or saline, polyalkylene glycols such as polyethylene glycol, oils of vegetable origin, or hydrogenated naphthalenes. Biocompatible, biodegradable lactide polymer, lactide/glycolide copolymer, or polyoxyethylene-polyoxypropylene copolymers may be used to control the release of the compounds. Other potentially useful parenteral delivery systems for SAP family member modulatory compounds include ethylene-vinyl acetate copolymer particles, osmotic pumps, implantable infusion systems, and liposomes. Formulations for inhalation may contain excipients, for example, lactose, or may be aqueous solutions containing, for example, polyoxyethylene-9-lauryl ether, glycocholate and deoxycholate, or may be oily solutions for administration in the form of nasal drops, or as a gel.

If desired, treatment with a SAP family member protein, gene, or modulatory compound may be combined with more traditional therapies for the disease involving excessive or insufficient levels of SH2 domain-containing protein-mediated signal transduction.

## VIII. Detection of Conditions Involving Altered SH2 Domain-Containing Protein-Mediated Signal Transduction

SAP family member polypeptides and nucleic acid sequences find diagnostic use in the detection or monitoring of conditions or diseases involving signal

-52-

transduction pathways which include SH2 domain-containing proteins. For example, increased expression of a SAP family member may be correlated with diseases hallmarked by decreased SH2 domain-containing protein-mediated signal transduction in humans. Accordingly, a decrease or increase in the level of SAP family member protein production may provide an indication of a deleterious condition. Levels of expression of SAP family member proteins may be assayed by any standard technique. For example, SAP family member expression in a biological sample (*e.g.*, a biopsy) may be monitored by standard Northern blot analysis or may be aided by PCR (see, *e.g.*, Ausubel *et al.*, *supra*; PCR Technology: Principles and Applications for DNA Amplification, H.A. Ehrlich, Ed. Stockton Press, NY; Yap *et al.*, Nucl. Acids. Res. 19: 4294, 1991).

Alternatively, a biological sample obtained from a patient may be analyzed for one or more mutations in SAP nucleic acid sequences using a mismatch detection approach. Generally, these techniques involve PCR amplification of nucleic acid from the patient sample, followed by identification of the mutation (*i.e.*, mismatch) by either altered hybridization, aberrant electrophoretic gel migration, binding or cleavage mediated by mismatch binding proteins, or direct nucleic acid sequencing. Any of these techniques may be used to facilitate mutant SAP family member detection, and each is well known in the art; examples of particular techniques are described, without limitation, in Orita *et al.* (Proc. Natl. Acad. Sci. USA 86: 2766-2770, 1989) and Sheffield *et al.* (Proc. Natl. Acad. Sci. USA 86: 232-236, 1989).

In yet another approach, immunoassays are used to detect or monitor SAP family member protein expression in a biological sample. SAP family member-specific polyclonal or monoclonal antibodies (produced as described above) may be used in any standard immunoassay format (*e.g.*, ELISA, Western blot, or RIA) to measure SAP family member polypeptide levels. These levels would be compared to wild-type SAP family member polypeptide levels. For example, an increase in SAP production may indicate a condition involving insufficient antigen-specific T cell activation. Examples of immunoassays are described, *e.g.*, in Ausubel *et al.*, *supra*.

-53-

Immunohistochemical techniques may also be utilized for SAP family member protein detection. For example, a tissue sample may be obtained from a patient, sectioned, and stained for the presence of SAP protein using an anti-SAP antibody and any standard detection system (e.g., one which includes a secondary antibody  
5 conjugated to horseradish peroxidase). General guidance regarding such techniques can be found in, e.g., Bancroft and Stevens (Theory and Practice of Histological Techniques, Churchill Livingstone, 1982) and Ausubel *et al.* (*supra*).

In one preferred example, a combined diagnostic method may be employed that begins with an evaluation of SAP family member protein production (for  
10 example, by immunological techniques or the protein truncation test (Hogerrorst *et al.*, Nature Genetics 10: 208-212, 1995) and also includes a nucleic acid-based detection technique designed to identify more subtle SAP family member mutations (for example, point mutations). As described above, a number of mismatch detection assays are available to those skilled in the art, and any preferred technique may be  
15 used. Mutations in SAP family member-encoding nucleic acids may be detected that either result in loss of expression of SAP family member or loss of normal SAP family member biological activity. In a variation of this combined diagnostic method, SAP family member biological activity is measured as antigen-specific T cell activation-inhibiting activity using any appropriate antigen-specific T cell activation  
20 assay system (for example, those described herein).

Mismatch detection assays also provide an opportunity to diagnose a SAP family member-mediated predisposition to diseases caused by inappropriate SH2 domain-containing protein-mediated signal transduction. For example, a patient heterozygous for a SAP family member mutation that induces a reduced level of SAP  
25 family member expression may show no clinical symptoms and yet possess a higher than normal probability of developing serious disease. Given this diagnosis, a patient may take precautions to minimize their exposure to adverse environmental factors (for example, UV exposure or chemical mutagens) and to carefully monitor their medical condition (for example, through frequent physical examinations). This type of SAP

-54-

family member diagnostic approach may also be used to detect SAP mutations in prenatal screens. The SAP family member diagnostic assays described above may be carried out using any biological sample (for example, any biopsy sample or other tissue) in which the SAP family member is normally expressed. Identification of a mutant SAP family member-encoding gene may also be assayed using these sources for test samples.

Alternatively, a SAP family member mutation, particularly as part of a diagnosis for predisposition to SAP family member-associated cancer, may be tested using a DNA sample from any cell, for example, by mismatch detection techniques. Preferably, the DNA sample is subjected to PCR amplification prior to analysis.

#### IX. Identification of Additional SAP Family Member Genes

Standard techniques, such as the polymerase chain reaction (PCR) and DNA hybridization, may be used to clone additional SAP family member homologues in other species. Southern blots of murine genomic DNA hybridized at low stringency with probes specific for human or murine SAP family member proteins reveal bands that correspond to SAP family members. Thus, additional SAP family member sequences may be readily identified using low stringency hybridization. Furthermore, region of high homology to a SAP family member may be used to screen databases for partial SAP family member sequences and SAP family member-specific primers may be used to clone additional SAP family member related genes by RT-PCR.

#### X. Construction of a Transgenic Animal

Characterization of the murine and human SAP family member genes (i.e., SAP protein encoding genes and EAT-2 protein encoding genes) provides information that is necessary for the construction of SAP family member knockout animal models by homologous recombination or by other standard techniques. Preferably, the model is a mammalian animal, most preferably a mouse. The murine SAP family member genomic DNA described herein enables the development of a mouse lacking an



-55-

endogenous SAP family member gene. Preferably, such a mouse with a SAP protein encoding gene knock-out may be used as a murine model of XLP disease.

Similarly, an animal model of SAP family member overproduction or an animal model of mutant SAP family member expression may be generated by  
5 integrating one or more SAP family member-encoding sequences into the genome operably linked to functional promoter sequences, according to standard transgenic techniques.

A replacement-type targeting vector, which would be used to create a knockout model, can be constructed using an isogenic genomic clone, for example,  
10 from a mouse strain such as 129/Sv (Stratagene Inc., LaJolla, CA). The targeting vector will be introduced into a suitably-derived line of embryonic stem (ES) cells by electroporation to generate ES cell lines that carry a profoundly truncated form of a SAP family member encoding gene. To generate chimeric founder mice, the targeted cell lines will be injected into a mouse blastula stage embryo. Heterozygous offspring  
15 will be interbred to homozygosity. Knockout mice would provide the means, *in vivo*, to screen for therapeutic compounds that modulate SH2 domain-containing protein-mediated signal transduction via a SAP family member-dependent pathway. Making such mice may require use of loxP sites due to the multiple copies of SAP family member encoding genes on the chromosome (see Sauer and Henderson, Nucleic Acids  
20 Res. 17: 147-61, 1989).

### EXAMPLES

The following examples are to illustrate the invention. They are not meant to limit the invention in any way.

#### Example I

#### 25 Cloning of SAP, a protein that interacts with the cytoplasmic domain of SLAM

To elucidate the signaling mechanism of the SLAM, co-activator proteins that interacted with the cytoplasmic domain of human SLAM were identified using a yeast

-56-

two-hybrid system with the 77 amino acid human SLAM cytoplasmic domain as "bait" (for a description, in general, of the yeast two-hybrid system, see, for example, Gyuris *et al.*, Cell 75:791-803, 1993). To do this, the cDNA encoding the human SLAM4 cytoplasmic domain (the sequence of which is shown on Fig. 1; SEQ ID NO: 1) was cloned in the pGBT9 vector (commercially available from Clontech Laboratories Inc., Palo Alto, CA) and used to screen a cDNA library derived from the human T cell line KT3 in the yeast two-hybrid system. The yeast two-hybrid assay and isolation of positive clones and subsequent interaction analyses are carried out as has been previously described (see, for example, PCT Publication WO 95/28497).

Eight human cDNA clones were isolated that encoded a 128 amino acid polypeptide chain, which was termed SLAM-Associated Protein (SAP; Fig. 2A and 2B). The 5' end of human SAP was obtained by the RACE technique (using reagents commercially available from Clontech). To confirm that full length cDNAs had been obtained, cDNA clones were also isolated from a human T cell library (Jurkat). We submitted the human SAP cDNA and protein sequence to the GenBank database, and it has been assigned GenBank Accession No. AF073019 (NID No. g3695070).

Having identified human SAP, we searched the databases and found a 200 nucleotide long fragment showing sequence showing identity to human SAP. Mouse SAP cDNA (Figs. 2C and 2D) was thus cloned by the screening of a mouse (C57BL6) thymus cDNA library in the Zap Express vector using as a probe a cDNA amplified from mouse thymic mRNA by RT-PCR (reverse transcriptase PCR), using specific primers flanking the 5' and 3' of the EST sequence AA255258. We submitted the murine SAP cDNA and protein sequence to the GenBank database, and it has been assigned GenBank Accession No. AF072903 (NID No. g3695066).

The predicted SAP protein contains a single SH2 domain (residues 6-102) followed in human SAP by a short 26 amino acid tail (residues 103-128 in human SAP) and in murine SAP by a short 24 amino acid tail (residues 103-126 in murine SAP). SAP was determined to contain an SH2 domain based on the close identity of five "blocks" of sequences within the putative SH2 domain with "blocks" found in

-57-

SH2 domain-containing proteins using the position-based method of Henikoff and Henikoff, J. Mol. Biol. 243: 574-578, 1994; Henikoff and Henikoff, Genomics 19: 97-107, 1994. Shown on Fig. 4A is the human SAP amino acid sequence with the positions of the five SH2 domain-hallmarking "blocks" shown in bold. In Fig. 4B, the sequences of these blocks is shown. Each of these "blocks" has a high degree (e.g., above 60%) sequence identity with a corresponding block in the SH2 domain of an SH2 domain-containing protein.

However, since SAP was isolated using a two-hybrid screen in yeast, which do not phosphorylate tyrosine residues, we do not define SAP as an SH2 domain-containing protein.

Human and murine SAP are highly homologous (96% identical) both in the SH2 and tail domains. In Fig. 5, human and murine SAP amino acid sequences were compared with the SH2-domains which were found most similar in a computer aided search, namely human SHIP, murine EAT-2 (GenBank Accession No. AF020263; Thompson *et al.*, Oncogene 13: 2649-2658, 1996), and human Abl. A computer-aided search for the SAP cDNA in the expressed-sequence tag library (EST) led to the identification of 3 human clones (Accession Nos. N89899, W19453 (Fetal Lung), and AA354319 (Jurkat)), and one highly homologous murine T cell derived clone of 294 bp (Accession No. AA255258). Whereas its SH2-domain was clearly related to other SH2-domains, particularly those of SHIP (45% identical), EAT-2 (40% identical), and Abl (35% identical), the short tail of SAP was unlike any known protein domain.

We next generated rabbit antisera containing polyclonal anti-SAP antibodies. To do this, human SAP (hSAP; CQGTTGIREDPDV) and murine SAP (mSAP; CQAPTGRRDSI) peptides were coupled to keyhole limpet hemocyanin (KLH; commercially available from Pierce Chemical Co., Rockford, IL) and used to inject New Zealand rabbits. Rabbits antisera were obtained using standard methods (see, for example, Ausubel *et al.*, Current Protocols in Molecular Biology, John Wiley & Sons, New York, NY, 1994).

The anti-human SAP antibodies were used to prepared immunoprecipitates of

-58-

whole lysates prepared from human T cell tumor cells (Jurkat) or human peripheral blood leukocytes (PBL) (Fig. 6A). Likewise, the anti-murine SAP antibodies were used to prepare immunoprecipitates of whole lysates prepared from murine (C57BL/6) thymocytes (Fig. 6B). The crude lysate was first precleared using 50  $\mu$ l of protein G-agarose beads (GIBCO BRL) and 5  $\mu$ l of normal mouse serum or normal rabbit serum for 1 hour. Immunoprecipitations were carried out using the indicated antibodies and 30  $\mu$ l of protein G-agarose beads for 3 hours at 4°C. Beads were then washed as described before (Ramesh *et al.*, Proc. Natl. Acad. Sci. USA 94: 14671-14676, 1997), and bead-associated proteins resolved by SDS-PAGE. The SDS-PAGE resolved proteins were transferred onto PVDF filters (commercially available from Millipore Corp., Bedford, MA), which were then blocked for 1 hour with 5% Skim-milk (or 3% Bovine Serum Albumin). Following Western blotting analysis with anti-human SAP (Fig. 6A) or anti-murine SAP (Fig. 6B), bound antibody was revealed using horseradish peroxidase-conjugated secondary antibodies using enhanced chemiluminescence (Supersignal; commercially available from Pierce Chemical Co.).

Antibodies directed at human SAP tail sequences detected a 15 kD protein in anti-SAP immunoprecipitates of detergent cell lysates made either from a human T cell tumor (Jurkat) or human peripheral blood leukocytes (PBL) (Fig. 6A). Likewise, antibodies directed at murine SAP tail sequences detected a 15 kD protein in detergent lysates made from murine thymocytes (Fig. 6B). Because the observed molecular weight was consistent with the predicted molecular mass, the human and mouse SAP cDNAs are unlikely to encode a fragment of a larger protein.

We next generated reagents to determine if SLAM and SAP interacted. cDNA encoding the entire coding region of human SAP (SEQ ID NO: 3) was next inserted into the GST fusion protein expression vector pGEX2T (commercially available from Pharmacia Biotech, Uppsala, Sweden) to generate the GST-SAP construct. The GST-SAP fusion protein-encoding plasmid was then used to transform *E. coli* bacteria, and, after propagation of the transformed bacteria, GST-SAP fusion

-59-

proteins were purified from lysed bacteria according to standard protocols. The SLAM construct was generated subcloning human SLAM4 cDNA in vector pJFE14-SR $\alpha$ . EL-4 cells were stably transfected by electroporation with a cDNA encoding SLAM4 or vector alone. Cells then were selected by growth in media with  
5 600 mg/ml neomycin for three weeks. Cells then were stained with anti-SLAM PE conjugated antibodies (A12) and positive cells were sorted.

Following cell surface biotinylation (using the EZ-Link Sulfo-NHS-Biotin reagent according to manufacturer's instructions; Pierce Chemical Co.) of untransfected EL-4 cells and SLAM transfected EL4 cells, whole lysates were  
10 prepared and incubated for 3 hours with 5  $\mu$ g of GST or SAP-GST fusion protein in the presence of glutathione beads. Following incubation, the beads were washed three times, resolved by SDS-PAGE, and transferred onto PVDF filters, which were then blocked for 1 hour with 5% Skim-milk (or 3% Bovine Serum Albumin). In Fig. 6C, Western blotting analysis was performed using streptavidin-HRP (horseradish  
15 peroxide-conjugated streptavidin, which specifically binds biotinylated proteins).

SLAM and SAP interact in T lymphocytes, as shown by the ability of a human SAP-GST fusion protein to specifically precipitate SLAM from detergent lysates made from a murine T cell (EL-4) transfected with human SLAM (Fig. 6C). In contrast, no interaction was detected in lysates made with untransfected EL4 cells  
20 (Fig. 6C). SAP, as detected by Western blotting with anti-SAP antibody, was also co-immunoprecipitated with anti-SLAM antibodies in detergent lysates prepared from PHA-activated human peripheral blood lymphocytes (as described in Isomaki *et al.*, J. Immunol. 159: 2986-2993, 1997) (top panel, Fig. 6D). Immunoprecipitations with anti-SLAM antibodies were performed as described above for anti-SAP  
25 immunoprecipitation.

### Example II

#### Identification of SAP as a T cell protein

We next did a series on Northern blotting analyses with SAP probes to

-60-

determine the tissue distribution of SAP mRNA. poly(A<sup>+</sup>) RNA from various human tissues (Clontech) or from total mRNA obtained from cell lines, according to standard techniques and using the Trizol reagent (commercially available from GIBCO BRL). Northern blotting analysis was performed (see, for example, Ausubel *et al.*, *supra*),  
5 using a radiolabelled human SAP-specific cDNA as a probe.

The level of human SAP mRNA expression was found to be highest in the thymus and lower in spleen and peripheral blood lymphocytes (Fig. 7A). SAP was expressed in all major subsets of human T cells (CD4<sup>+</sup>, CD45RO<sup>+</sup>, CD45RA<sup>+</sup>, and CD8<sup>+</sup>) (Fig. 7C) and in the T cell tumor, Jurkat, and the EBV<sup>+</sup> Burkitt lymphoma  
10 line, Raji. No transcripts were detected in the EBV<sup>+</sup> Burkitt lymphomas, Namalwa and BL41/B95-8; the EBV transformed cell line, X50-7; the EBV<sup>+</sup> marmoset cell lines, B95-8 and FF41; the EBV<sup>-</sup> Burkitt lymphoma, BJAB; or the pre-B ALL-lines, LAZ 221 and NALM 6 (Figs. 7B and 7C, and data not shown). The low level of SAP mRNA that was detected in the small intestine probably resulted from lymphocyte  
15 contamination. In *wt* mice, SAP is expressed in T cells, but not in B cells, while no SAP mRNA or protein could be detected in CD3 $\epsilon$ <sup>Null</sup> mice which lack T cells. Both human mRNA species encoding SAP (2.5 and 0.9 kb) were represented amongst the cDNA clones isolated in the two-hybrid system and encoded the same open reading frame, but differed in their 3' untranslated sequences. Collectively, these studies  
20 showed that, unlike SLAM, SAP was primarily expressed in T cells.

### Example III

#### SAP is encoded by the X-linked Lymphoproliferative disease (XLP) gene

Using a 45 kb pBAC clone that contained all four exons of murine SAP (see Fig. 5), the SAP gene was localized within band A5.1 of the murine X chromosome.  
25 Because of synteny between murine band A5.1 and human Xq25, the locus at which the immune deficiency X-linked Lymphoproliferative disease (XLP) had been mapped (Lamartine *et al.*, Eur. J. Hum. Genet. 4: 342-351, 1996; Porta *et al.*, Genome Research. 7: 27-36, 1997; Lanyi *et al.*, *supra*; Purtilo, D.T., *supra*), it was plausible

that the SAP gene was involved in this disease. Moreover, the observation that uncontrolled B cell proliferation following EBV infection in XLP patients often leads to fatal infectious mononucleosis, malignant lymphomas, hypogammaglobinemia, or aplastic anemia (Puttilo, D.T., *supra*; Seemayer *et al.*, *Pediatr. Res.* 38: 471-478, 1995) was in agreement with the involvement of SLAM in T $\leftrightarrow$ B cell interactions. To test the hypothesis that SAP was encoded by the XLP gene, SAP cDNAs were isolated from peripheral blood mononuclear cells of three XLP patients: a previously described patient A1 (Rousset *et al.*, *Clin. Exp. Immunol.* 63: 280-289, 1986), and a set of two brothers, B1 and B2, who were recently diagnosed with XLP.

XLP patient A1 developed hypogammaglobunemia and recurrent pulmonary infections a few months after severe EBV induced infectious mononucleosis (Rousset *et al.*, *supra*). His family history and the persistence of an unbalanced virus-host relationship after EBV infection are consistent with the main characteristics of XLP syndrome (Rousset *et al.*, *supra*). We used RT-PCR (reverse transcriptase-PCR) to elucidate whether or not patient A1 had a mutation in the SAP gene. RT-PCR was performed using GenAmp RT-PCR kit (Perkin Elmer Corp., Norwalk, CT) using the following primers: forward primer 5'-GCC TGG CTG CAG TAGCAG CGG CAT CTC CC-3' (SEQ ID NO: 17); and reverse primer 5'-ATG TAC AAA AGT CCA TTT CAG CTT TGA C-3' (SEQ ID NO: 18). The RT-PCR products were then subjected to 10% polyacrylamide gel electrophoresis at 200 volts for 7 hr. The gels were stained with CYBR Green (Molecular Probes Inc., Eugene, OR) for 20 min., according to manufacturer's procedures, and visualized and photographed using a UV illuminator.

When RT-PCR products of T cells from patient A1 were analyzed by polyacrylamide gel electrophoresis (Fig. 8A), initially three products were found with the sizes: 629 bp (the full length SAP coding sequence); 565 bp; and 520 bp. Upon cloning of the RT-PCR products of patient A1 four different mRNAs were identified: (i) full length hSAP; (ii) a coding sequence,  $\Delta$ E2, lacking the 64 nucleotides of exon 2 (A1-1 in Figs. 3, 9A, and 9B); (iii) E3 $\Delta$ 55 with a 55 nucleotide deletion in exon 3

-62-

(hSAP $\Delta$ 55 in Figs. 3, 9A, and 9B); and (iv) a cDNA with both a deleted exon 2 and the deletion E3 $\Delta$ 55 (A1-2 in Figs. 3, 9A, and 9B). Thus, T cells from patient A1 had four mRNA species coding for different forms of SAP, two of which were detected in healthy individuals (629 and 574 bp) and two that were specific for the patient (A1-1 of 565 bp and A1-2 of 520 bp). (Note that the 574 and 565 bp species cannot be separated on PAGE in Fig. 8A.) The predicted amino acid sequences suggested two truncated proteins in which essentially only the first exon of SAP was intact.

RT PCR products from peripheral blood mononuclear cells of 60 healthy individuals contained two mRNA species (629 and 574 bp), as judged by polyacrylamide gel electrophoresis and nucleotide sequence analyses (Figs. 8A and 9B). These represented full length hSAP and E3 $\Delta$ 55 (hSAP $\Delta$ 55 in Figs. 3, 9A and 9B), in which part of exon 3 had been deleted. The E3 $\Delta$ 55 coding sequence started at the beginning of exon 3 at nucleotide position 288 and ended at nucleotide 342 (Fig. 9A). Because of a frame shift, the predicted protein sequence was identical to that of the first two exons of SAP (amino acids 1-67) followed by a nine amino acid sequence and a stop codon (Fig. 9B). Taken together, the mRNAs of patient A1 either had a deletion of exon 2, or a deletion of exon 2 in addition to a deletion of exon 3, which was a normal variant. These mutant forms derived from patient A1 were estimated to represent 90% of the patient's SAP mRNA.

To aid our analysis, each of the four exons of human SAP gene was amplified by PCR using the following primers:

forward primer 5'-GCC CTA CGT AGT GGG TCC ACA TAC CAA CAG-3' (SEQ ID NO: 19), and reverse primer 5'-GCA GGA GGC CCA GGG AAT GAA ATC CCC AGC-3' (SEQ ID NO: 20) for exon 1;

forward primer 5'-GGA AAC TGT GGT TGG GCA GAT ACA ATA TGG-3' (SEQ ID NO: 21), and reverse primer 5'-GGC TAA ACA GGA CTG GGA CCA AAA TTC TC-3' (SEQ ID NO: 22) for exon 2;

forward primer 5'-GCT CCT CTT GCA GGG AAA TTC AGC CAA CC-3' (SEQ ID NO: 23), and reverse primer 5'-GCT ACC TCT CAT TTG ACT TGC TGG CTA



-63-

CAT C-3' (SEQ ID NO: 24) for exon 2; and forward primer 5'-gac agg gac cta ggc tca ggc ata aac tga c-3' (SEQ ID NO: 25), and reverse primer 5'- ATG TAC AAA AGT CCA TTT CAG CTT TGA C -3' (SEQ ID NO: 26) for exon 4. These PCR primers were designed based on the genomic  
5 sequence of human SAP released by the EMBL database (Accession No. AL022718). The PCR products were visualized on 2% agarose gels, and subcloned into pCR2.1 vector (commercially available from Invitrogen Corp., Carlsbad, CA followed by sequencing.

The 55 nucleotide deletion in exon 3 in healthy individuals was readily  
10 explained by a infrequently used splicing acceptor site within that exon: CAG preceded by a CT rich stretch of nucleotides (Lewin, B., Genes V, pp. 1-1272, Oxford University Press, Oxford, 1994) (see Fig. 10A). However, in order to explain the frequent deletion of exon 2 in patient A1, the intron sequences surrounding exon 1, 2, and 3 needed to be determined. To this end, a PCR assay based upon sequences of at  
15 least 150 nucleotides of all four introns was designed. This was facilitated by a very recent sequence in the EMBL database (Accession No. AL022718) stating that the XLP gene encoded a protein, termed DSHP. Its cDNA, genomic DNA, and protein sequences were identical to human SAP and, in addition, the exon/intron boundaries reported were identical to those determined for the mouse gene (see Fig. 5). Upon  
20 sequencing of the 443 bp PCR product derived from the exon 2 area of patient A1's DNA, a C=G mutation was found in the nucleotide adjacent to the exon 2 splice acceptor site in the DNA of patient A1 (Fig. 10A). This mutation in other gene sequences is known to affect the efficiency of the splice acceptor site (Lewin, B., *supra*) and explained the skipping of exon 2 in the majority (~ 90%) of the patient's  
25 SAP mRNA species.

To exclude the possibility that the point mutation found in XLP patient A1 represented a genetic polymorphism, a restriction enzyme assay was designed. This assay was based on the fact that the C=G mutation generated a novel Mnl I restriction enzyme recognition site in the 443 bp exon 2 PCR product. For the analysis of a

-64-

point mutation in the region preceding exon 2, exon 2 PCR products were gel-purified (using reagents commercially available from Qiagen Inc., Santa Clarita, CA) and digested with restriction enzyme Mnl I (commercially available from New England Biolabs Inc., Beverly, MA), followed by 10% polyacrylamide gel electrophoresis at 200 volts for 3 hr. As shown in Fig. 10B, digestion with Mnl I generated four fragments (187, 151, 64, and 41 bp) with A1's 443 bp PCR product, while DNA samples from 108 healthy individuals generated three fragments (251, 151, and 41 bp) (see, *e.g.*, CT-1, CT-2, and B3 in Fig. 10B). Because patient A1 was Italian, DNA from 50 healthy Italian females was analyzed as controls (providing 100 X-chromosomes). An additional 86 X-chromosomes were from 28 female (*i.e.*, 56 X-chromosomes) and 30 male (*i.e.*, 30 X-chromosomes) individuals with a random genetic background from area of the Boston, Massachusetts. Collectively, the data demonstrate that the XLP patient A1 had a mutation affecting the exon 2 splice acceptor site, which gave rise to severely truncated forms of SAP. Since this mutation was not detected in 186 X-chromosomes from healthy individuals, it did not represent a genetic polymorphism.

Two additional XLP patients, brothers B1 and B2, had a deletion of the SAP gene, while a healthy brother B3 had a normal SAP gene. The first patient, B1, was a 23 year old male with a history of recurrent pulmonary infections, dysgammaglobulinemia (elevated IgA and IgM), poor specific antibody responses to tetanus toxoid antigen, and depressed T cell proliferative response to mitogens. He developed fever, pneumonia, and hilar adenopathy, which quickly progressed to fulminant infectious mononucleosis with hemophagocytosis. EBV infection was documented by PCR and serology. One of B1's male siblings, B2, also suffered from the XLP syndrome, which manifests itself as pancytopenia, splenomegaly, dysgammaglobulinemia, and depressed T cell proliferative responses to mitogens. A third brother, B3, appears healthy, and two additional brothers, of whom no material was available, had classical XLP symptoms: one had recurrent EBV+ non-Hodgkin's lymphomas, and another died with the diagnosis of Wegener's granulomatosis and

-65-

pulmonary infiltrates.

In contrast to the healthy sibling B3, no RT-PCR product encoding hSAP could be isolated from the peripheral blood mononuclear cells of patients B1 and B2 (Fig. 8A). This was because no mRNA could be detected in Northern blotting experiments with RNA from patients B1 (not shown) and B2 (Fig. 8B). This result suggested that either the SAP gene was not transcribed in T cells from B1 and B2, or that the gene was deleted. Since none of the four exons encoding SAP could be generated in a PCR analysis based upon genomic DNA from B1 and B2 (Fig. 10C), and because a control PCR reaction for exon 2 of BRCA1 gave the expected product, we conclude that the SAP gene was deleted in XLP patients B1 and B2. By contrast, DNA from the healthy brother B3 generated PCR fragments of the expected size and with the wt sequence. Taken together, the analyses of XLP patients A1, B1, and B2 demonstrated the XLP gene encodes SAP.

#### Example IV

#### 15 The SH2-domain of SAP binds to a specific sequence in the cytoplasmic domain of SLAM

Identification of SAP as the gene product altered in XLP raised the question of how this SLAM interactive T cell protein could account for the immunologic disturbances associated with disease. The SLAM cytoplasmic domain contains three Tyr residues (Y281, Y307, and Y327) that are surrounded by consensus SH2-domain-binding-sequences (Cocks *et al.*, *supra*), suggesting the possibility of an interaction with the SH2-domain of SAP. However, SAP was found to bound to SLAM in a yeast two-hybrid system, a system in which tyrosine residues are not phosphorylated. Hence, the SAP-SH2 domain had to bind to SLAM in a more complex fashion than through a classical interaction with a short phospho-tyrosine containing peptide.

In order to determine the portion of the cytoplasmic domain of SLAM to which the SAP SH2-domain bound, constructs encoding the CD8 extracellular and transmembrane domains, and truncated cytoplasmic domains of SLAM were

-66-

generated, and schematic diagrams of which are shown on Fig. 11A. Shown in Fig. 11B are the amino acid sequences of the cytoplasmic domains of SLAM3 and SLAM4 that were fused to CD8 extracellular and transmembrane domains. In addition, a truncation of the cytoplasmic domain of SLAM4, the sequence of which is shown in Fig. 11B (SLAM 4 del 1), was generated by PCR from a full-length SLAM4 template, and the PCR product used to generate the CD8-SLAM del 1 fusion protein. The CD8-SLAM constructs were cloned in pCDNA3 (commercially available from Invitrogen Corp.). Human SAP cDNA was cloned in expression vector pCMV2-FLAG (commercially available from KODAK), thus generating a Flag-tagged version of human SAP. The CD8-SLAM constructs were co-expressed with Flag-tagged human SAP in COS-7 cells by transforming the cells using the DEAE-dextran method (Ausubel *et al.*, *supra*).

Following cell surface biotinylation using the EZ-Link Sulfo-NHS-Biotin reagent according to manufacturer's instructions (Pierce Chemical Co.), cell lysates which were prepared, clarified by centrifugation at 14,000 x g for 15 min. at 4°C, and the crude lysate precleared using 50 µl of protein G-agarose beads (GIBCO BRL, Gaithersburg, MD) and 5 µl of normal mouse serum or normal rabbit serum for 1 hour. Anti-CD8 (OKT3) and anti-SLAM immunoprecipitations were next performed using the antibodies and 30 µl of protein G-agarose beads for 3 hours at 4°C. The beads were then washed as described (Ramesh *et al.*, *supra*). Crude lysates and immunoprecipitates were resolved by SDS-PAGE, and transferred onto PVDF filters, which were then blocked for 1 hour with 5% Skim-milk (or 3% Bovine Serum Albumin), and then probed with anti-SAP antibodies (top panel, Fig. 12A) or streptavidin (bottom panel, Fig. 12A). Bound antibody was revealed using horseradish peroxidase-conjugated secondary antibodies using enhanced chemiluminescence (Supersignal; commercially available from Pierce Chemical Co.).

Both CD8-SLAM3 (derived from a natural variant of SLAM having a short cytoplasmic domain containing only Y281) and CD8-SLAM del1 (containing Y281 and Y307), were able to co-precipitate SAP with the same efficiency as SLAM itself

-67-

(Fig. 12A). In contrast, two control chimeric proteins comprising the CD8 extracellular and transmembrane domains, and either the CD3- $\epsilon$  or the CD3- $\zeta$  cytoplasmic domain, did not co-precipitate SAP (Fig. 12A). This suggested the presence of a specific SAP binding site around the most membrane proximal tyrosine residue (Y281) of the SLAM cytoplasmic tail.

We next generated the following peptides:

SLAM Y1: CVEKKSLTIYAQVQK (SEQ ID NO: 27); SLAM pY1: CVEKKSLTIpYAQVQK (SEQ ID NO: 28); SLAM F1: CVEKKSLTIFAQVQK (SEQ ID NO: 29); SLAM Y2: CTTIYVAATEPVPESVQE (SEQ ID NO: 30); and  
SLAM Y3: CTVYASVTLPES (SEQ ID NO: 31). (Note: Rabbit polyclonal antibodies directed toward SLAM Y2 and SLAM Y3 peptides were generated as described above.) Following synthesis, the peptides were coupled to beads (sulfolink coupling gel, Pierce Chemical Co.). Coupled peptides were incubated for 1 hour with lysates from mouse thymocytes or from CD8SLAM3 transiently transfected cells.  
Where indicated in Figs. 12B and 12C, the coupled peptides were incubated with lysate in the presence of different concentrations of soluble peptides. Following incubation, the beads were washed three times and bead-bound proteins resolved by SDS-PAGE.

As predicted from the *in vivo* experiments, only a peptide (SLAM Y1) containing Y281, coupled to agarose beads, was able to specifically precipitate SAP from murine thymocyte lysates, but not peptides containing the other tyrosines of the cytoplasmic domain of SLAM (Fig. 12B). More importantly, binding of SAP to the agarose-coupled SLAM Y1 was blocked only by the same peptide regardless whether Y281 was replaced by Phe (SLAM F1) or by a peptide in which phospho-Tyr replaced Y281 (SLAM pY1) (Fig. 12C). This confined the binding site to a short amino acid sequence in SLAM and confirmed that phosphorylation of Y281 was not required for interactions between the two polypeptide chains. However, the SH2-domain of SAP was functionally intact (*i.e.*, it could still bind phosphorylated tyrosine residues) since it could be purified on a Sepharose-phospho-Tyr column (data not

-68-

shown).

Human SAP cDNA and mutant SAP cDNAs, generated by PCR (SAP del1, SAP del2, SAP del3, SAP del4; see Fig. 5), were next cloned in expression vector pCMV2-FLAG (commercially available from KODAK). These four SAP mutants  
5 with deletions of the SAP-tail, when co-transfected into COS-7 cells with the CD8-SLAM3 chimera, were able to interact with the short cytoplasmic domain of SLAM3, which contained the Y281 segment, as determined following anti-CD8 immunoprecipitation and Western blotting analysis with anti-FLAG antibody (Fig. 12D). Collectively these data demonstrated conclusively that SLAM interacted with  
10 SAP via part of its SH2-domain.

#### Example V

#### SAP blocks recruitment of the tyrosine phosphatase SHP-2 to the phosphorylated cytoplasmic domain of SLAM

The novelty of SLAM-SAP recognition, wherein the usual requirements for  
15 SH2-domain interactions appeared to be altered, raised the possibility that SAP was blocking physiological relevant interactions in the function of SLAM, thus acting as a natural inhibitor molecule. This model also predicted that SAP mutants that were unable to bind to SLAM would not act as inhibitors. This hypothesis was testable, because we have observed that upon phosphorylation by *c-fyn*, SLAM did recruit the  
20 tyrosine-phosphatase SHP-2.

First, two SAP mutants were expressed in COS cells: the A1 deletion ( $\Delta$ exon2+E3 $\Delta$ 55) and a classical mutation in the SH2 domain which affects binding of phosphotyrosine, SAP 32R $\rightarrow$ Q (*i.e.*, the R residue at position 32 changed to a Q residue). Neither mutant bound to the cytoplasmic tail of SLAM as judged by *in vivo*  
25 (Fig. 13A) or *in vitro* (Fig. 13B) assays. Thus, neither SAP-mutant could act as a dominant negative mutant via SLAM. More importantly, the experiment with mutant SAP 32R $\rightarrow$ Q showed that the interaction of SAP with SLAM depended upon an intact phospho-tyrosine binding pocket. Interestingly, an XLP patient with a mutation in

-69-

SAP R32 has recently been found. Taken together, the absence of binding of the two SAP mutants found in patients further supports the model that absence of SAP/SLAM interactions are critical for this disease.

Next, SLAM and *c-fyn* were co-transfected with or without SAP into COS-7  
5 cells, which contain endogenous SHP-2. If SLAM was phosphorylated by *c-fyn*, SHP-2 was co-precipitated with anti-SLAM (Fig. 13C), but in the absence of *c-fyn* SLAM did not bind SHP-2. Introduction of SAP, together with SLAM and *c-fyn*, completely blocked the interaction of SHP-2 and SLAM. The absence of SHP-2 in the presence of SAP could not be due to lower levels of phosphorylation of SLAM,  
10 because a significantly larger percentage of SLAM molecules was phosphorylated in the presence of SAP than in its absence. These results were confirmed in reciprocal co-immunoprecipitation studies with anti-SAP reagents. By contrast, as shown in Fig. 13D, mutant SAP 32R→Q did not block the binding of SHP-2 to SLAM. A similar result was obtained with the A1 deletion mutant. Thus, recruitment of SHP-2  
15 to phosphorylated SLAM was blocked by the binding of SAP to the sequence surrounding the most membrane proximal tyrosine residue Y281 of SLAM.

Since SHP-2 itself can act as a negative regulator of signal transduction cascades (Marengere *et al.*, *supra*), binding of SAP could have a positive effect on co-stimulation by anti-SLAM. To test this, SAP was over-expressed by transient  
20 transfections into the Jurkat T cells, together with an IL-2-promoter-luciferase reporter construct and SLAM. cDNAs coding for SLAM and SAP were transiently transfected with an IL-2 promoter-luciferase constructs into Jurkat-Tag cells, as described (Martinez-Martinez, S. *et al.*, Mol Cell Biol. 17: 6437-6447, 1997). After 24 hours, transfected cells were stimulated with anti-CD3 and/or anti-SLAM  
25 monoclonal antibodies, as described (Cocks *et al.*, *supra*). Postnuclear lysates were analyzed in a Berthold luminometer 8 hours later.

Stimulation of SLAM+SAP transfected cells with a combination of anti-SLAM and anti-CD3 monoclonal antibodies resulted in a significantly higher IL-2-promoter-luciferase reporter activation, than that observed in Jurkat cells transfected

-70-

with SLAM alone (with an IL-2-promoter-luciferase reporter construct) (Fig. 14). Anti-SLAM by itself did not induce the IL-2-promoter-luciferase reporter signal, and the signal induced by anti-CD3 reagents was lower than that generated by the two antibodies together. Thus, over-expression of SAP in T cells had a positive effect on  
5 the co-stimulatory activity of anti-SLAM.

#### EXAMPLE VI

##### Mapping the SAP gene

Given our finding that the SAP gene mapped to the human XLP locus, we next produced a finer map of both the human and the murine SAP genes. This finer  
10 mapping allows the identification of putative transcription factors that may bind to the SAP gene and, thus, control SAP protein expression.

On Figs. 15-18 are shown the DNA sequences of the 5' region / exon1 / intron1 of the murine SAP gene (SEQ ID NO: 14), the DNA sequences of exon2 / intron2 of the murine SAP gene (SEQ ID NO: 15), and the DNA sequences of  
15 intron2 / exon3 / intron3 / exon4 / intron4 of the murine SAP gene (SEQ ID NO: 16), respectively. From these sequences, we constructed a schematic diagram showing the locations of the four exons of the murine SAP gene on the X chromosome. This information allows the production of a SAP knock-out animal, or a transgenic animal expressing a mutant or truncated SAP protein (see above). Methods for generating  
20 such animals are well known (see for example, Fung-Leung *et al.*, Cell 65:443-9, 1991; Kozieradzki *et al.*, J. Immunol. 158:3130-3139, 1997; Galli-Taliadoros *et al.*, J. Immunol. Methods 181:1-15, 1995).

In addition, as shown on Fig. 19, we identified a number of putative binding sites for transcription factors in the non-coding regions of both human and murine  
25 SAP (*i.e.*, the 5' untranslated regions and the introns) which may regulate the expression of SAP. By specifically disrupting one or more of these binding sites, the level of expression of SAP may be altered in a patient suffering from a disease



-71-

characterized by an inappropriate amount of SH2-domain containing protein mediated signal transduction.

#### EXAMPLE VII

##### Higher Expression of SAP in Th1 than Th2 helper T cells

- 5           Because engagement of SLAM during antigen-specific T-cell stimulation has been shown to induce IFN- $\gamma$  production and to redirect the Th2 phenotype to a Th1/Th0 phenotype (Carballido *et al.*, J. Immunol. 159: 4316-4321, 1997), we next sought to determine if Th1 and Th2 helper T cells had different levels of expression of SAP. Accordingly, we isolated Th1 and Th2 cells from mice according to standard
- 10 techniques. Briefly, CD4<sup>+</sup> T cells from mice expressing the transgene for the DO11.10  $\alpha\beta$ -TCR, which recognizes residues 323-339 of chicken ovalbumin (OVA) in association with I-A<sup>d</sup>, were cultured in complete RPMI 1640 media with OVA 323-339 peptide (1  $\mu$ M) and mitomycin-treated splenocytes. For Th1 phenotype T cell development, recombinant murine IL-12 (10 ng/ml), neutralizing anti-IL-4
- 15 monoclonal antibody (clone 11B11, 40  $\mu$ g/ml, commercially available from R&D Ssystems, Minneapolis, MN) was added. For Th2 phenotype T cell development, recombinant murine IL-4 (10 ng/ml) and neutralizing polyclonal anti-murine IL-12 (clone TOSH-2, 3  $\mu$ g/ml, commercially available from Endogen, Cambridge, MA) was added.
- 20           Cells were cultured for three rounds of antigenic stimulations under polarizing conditions, and then stimulated on plate-bound anti-CD3 (clone 2C11, commercially available from PharMingen, San Diego, CA) in the presence of human IL-2 (commercially available from Endogen) for 6 hours. After no stimulus or stimulus for 6 hours with, polyA<sup>+</sup> RNA was prepared from the cells (using the FastTrack mRNA
- 25 kit commercially available from Invitrogen, Carlsbad, CA), electrophoretically resolved, and subjected to Northern blotting analyses using as probes the following radiolabelled murine DNAs: Interferon-gamma (IFN- $\gamma$ ; to identify Th1 cells), interleukin-4 (IL-4; to identify Th2 cells), mSAP (which identifies a 0.9 kB band),

-72-

mSLAM (which identifies a 2.1 kB band) and  $\beta$ -actin (control; identifies a 2.2 kB band).

As shown on Fig. 20, following 6 hours of stimulation, the 2.1 kB SLAM mRNA band increased in both stimulated (vs. non-stimulated) Th1 and Th2 cells.

- 5 The 0.9 kB SAP mRNA, however decreased in both Th1 and Th2 cells following 6 hours of stimulation (note that the decrease was less noticeable in the Th1 cell population). However, it is clear that in non-stimulated cells, the level of SAP mRNA species was higher in Th1 cells than in Th2 cells. Given this finding, in order to shift the immune response to a Th1-mediated response as opposed to a Th2-mediated
- 10 response (e.g., in the treatment of leprosy), it is advisable to treat the patient with a compound that causes an increase in the level of expression of the SAP mRNA in resting T cells.

### EXAMPLE VIII

#### Identification of EAT-2 as a SAP family member protein

- 15 The murine EAT-2 protein was identified as an SH2 aberrantly expressed in Ewing's sarcoma tumor cell lines (Thompson *et al.*, Oncogene 13:2649-58, 1996; GenBank Accession Nos. AF020264 (human, fragment) and AF020263 (mouse)). Given our finding of the high sequence identity between the SH2 domain of EAT-2 with SAP, as well as the similarity in the overall length of the proteins, we sought to
- 20 determine if EAT-2 was a second member of the SAP family of proteins.

- To do this, we subcloned the entire coding region of the EAT-2 cDNA into a FLAG-fusion expression vector cloned in expression vector pCMV2-FLAG (commercially available from KODAK) to create a EAT2-FLAG fusion protein terminally fused to the FLAG tag. As in Example IV, the EAT2-FLAG encoding
- 25 expression vector was co-transfected with the CD8SLAM3 encoding vector into COS cells. Following cell surface biotinylation using the EZ-Link Sulfo-NHS-Biotin reagent (Pierce Chemical Co.), the COS cells were lysed and immunoprecipitated with anti-CD8 antibody. The immunoprecipitates were resolved by SDS-Page and

-73-

subjected to Western blotting analyses with anti-FLAG antibody, anti-SAP antibody, or streptavidin. As can be seen on Fig. 21, both Flag tagged SAP and Flag tagged EAT2 can be pulled down with CD8SLAM3 (Fig. 21, top panel). Thus, EAT2, like SAP, binds the cytoplasmic tail of SLAM3, which bears only the one tyrosine residues, Y281. Equivalent loading of the lanes is shown at the bottom panel of Fig. 21, with equivalent binding of streptavidin to all three lanes.

### EXAMPLE IX

#### Isolation of SAP associated proteins by yeast 2-hybrid analysis

A plasmid which encodes the GAL4 DNA-binding domain fused to the entire coding domain of human or murine SAP, is constructed by inserting DNA encoding the coding region of human or SAP into a GAL4 DNA binding domain fusion protein expression vector pGBT9 (Clontech, Palo Alto, CA). GAL4 DNA binding domain-SAP fusion protein is then used as "bait" (DNA-binding domain hybrid) in yeast two-hybrid screens of a human cDNA "prey" library of PHA-stimulated peripheral blood leukocytes (PBL) or a murine cDNA "prey" library of a T cell lymphoma (both commercially available from Clontech). The yeast two-hybrid assay and isolation of positive clones and subsequent interaction analyses are carried out as has been previously described (see, for example, PCT Publication WO 95/28497).

DNA sequencing of positive SAP-interacting clones is performed on an Applied Biosystems model 373A automated DNA sequencer. Full length cDNA encoding polypeptide fragments of SAP-interacting proteins are isolated by using the cDNA clone encoding the SAP-interacting polypeptide fragment to screen other libraries (e.g., longer cDNA libraries or genomic libraries) using standard hybridization techniques (see Ausubel *et al.*, *supra*; Sambrook *et al.*, *supra*).

### EXAMPLE X

#### Additional SH2 Domain-Containing Protein-Mediated Signal Transduction Assays

Numerous assays are known which evaluate SH2 domain-containing protein

-74-

mediated signal transduction. Such signal transduction pathways occur in, for example, epidermal cells through the Epidermal Growth Factor (EGF) receptor (Vogel *et al.*, Science 259: 1611-1614, 1993), fibroblasts through the Platelet Derived Growth Factor (PDGF) receptor (Fantl *et al.*, Cell 69: 413-423, 1992), and B cells (Ono *et al.*, Cell 90: 293-301, 1997).

In one particular example, SAP activity on antigen-specific T cell activation can be assessed using a variety of antigen-specific T cell activation assays are known in the art. Most are based upon the increased production of activation dependent T cell genes including, without limitation, the following cell surface molecules: Fas ligand, MHC class II, CD25, and CD69; as well as the following cytokines: interleukin-2 (IL-2), and  $\gamma$ -interferon. All of the following examples may be easily modified to identified compounds which affect SAP expression and biological activities, as well as peptide and non-peptide SAP mimetics which have SAP biological activity.

For example, a T cell specific for, for example, a viral peptide, may be stimulated with a syngeneic antigen presenting cell presenting that viral peptide on its cell surfaces in context with MHC. An activated T cell will express Fas ligand, MHC class II, CD25, and CD69 on its cell surface (as opposed to a resting, unactivated T cell which expresses none of these cell surface molecules). Similarly, an activated T cell, unlike a resting T cell, will produce IL-2 and  $\gamma$ -IFN. The cell surface activation markers can be detected by antibodies (commercially available from, for example, Becton Dickinson, San Jose, CA). Binding of these antibodies may be rapidly assessed by binding of a secondary labeled antibody, followed by detection of that bound secondary antibody (by, for example, FACS analysis or addition of colorimetric substrate for the label). The cytokines released by activated T cells may be detected from the supernatants of cells by commercially available ELISA kits (commercially available from, for example, Endogen Inc., Woburn, MA).

Antigen-specific T cell activation may also be determined by the production of activation-dependent cytokines, such as IL-2, in a proliferation assay of IL-2

-75-

dependent cells. For example, Jurkat human T cells may be stimulated with anti-CD3 plus anti-CD28 antibodies (commercially available from, for example, Pharmingen, San Diego, CA). Following culture for 24-48 hours at 37°C, the supernatants from these cells is collected, freeze-thawed, and added to an IL-2 dependent cell line, such as CTLL-20 (available from the ATCC). For proliferation studies, the CTLL-20 cells are then returned to culture for 48-72 hours total, with the addition of <sup>3</sup>H-thymidine to the cells in the final 6-20 hours of culture. The cells are then lysed, and proliferation was measured by amount of <sup>3</sup>H-thymidine uptake. Non-radioactive cell proliferation assays are also known in the art (*e.g.*, the CellTiter 96® AQueous Non-Radioactive Cell Proliferation Assay commercially available from Promega Co., Madison, WI).

Other examples of antigen-specific T cell activation assays are also provided in the following references: increased interleukin-2 (IL-2) production (Aversa *et al.*, J. Immunol. 158: 4036-4044, 1997); increased interferon-γ (IFN-γ) production (Aversa *et al.*, J. Immunol. 158: 4036-4044, 1997); increased CD69 expression (Zubiaur *et al.*, J. Immunology 159: 193-205, 1997); increased Fas Ligand expression (Ju *et al.*, Nature 373:444-448, 1995); increased interleukin-4 production (Cocks *et al.*, *supra*); increased incidence of phosphotyrosine-containing proteins (Zubiaur *et al.*, *supra*); and recruitment of phosphatidylinositol 3'-kinase (PI3-kinase) to the T cell receptor signaling complex (de Aoz *et al.*, J. Biol. Chem. 272: 25310-25318, 1997).

20

#### Other Embodiments

In other embodiments, the invention includes any protein which is substantially identical to the human SAP polypeptide provided in Fig. 2B (SEQ ID NO: 4), murine SAP polypeptide provided in Fig. 2D (SEQ ID NO: 6), or to GenBank Accession Nos. AF020263 and AF020264; such homologues include other substantially pure naturally-occurring mammalian SAP family member proteins as well as splice variants, allelic variants; natural mutants; induced mutants; DNA sequences which encode proteins and also hybridize to the SAP family member encoding DNA sequence of Fig. 2A (SEQ ID NO: 3), Fig. 2C (SEQ ID NO: 5),

-76-

GenBank Accession No. AF020263, or GenBank Accession No. AF020264, under high stringency conditions (*e.g.*, hybridizing at 2X SSC at 40°C with a probe length of at least 40 nucleotides) or, less preferably, under low stringency conditions (*e.g.*, hybridizing at 5X SSC at 25°C with a probe length of at least 80 nucleotides); and

5 proteins specifically bound by antisera directed to a SAP family member polypeptide. The term also includes chimeric polypeptides that include a portion derived from a SAP family member polypeptide.

The invention further includes analogs of any naturally-occurring SAP family member polypeptides. Analogs can differ from the naturally-occurring SAP family

10 member proteins by amino acid sequence differences, by post-translational modifications, or by both. Analogs of the invention will generally exhibit at least 50%, more preferably 70%, and most preferably 90% or even 95% identity with all or part of a naturally-occurring SAP family member amino acid sequence. The length of sequence comparison is at least 15 amino acid residues, preferably at least 25 amino

15 acid residues, and more preferably more than 35 amino acid residues. Modifications include *in vivo* and *in vitro* chemical derivatization of polypeptides, *e.g.*, acetylation, carboxylation, phosphorylation, or glycosylation; such modifications may occur during polypeptide synthesis or processing or following treatment with isolated modifying enzymes. Analogs can also differ from the naturally-occurring SAP family

20 member polypeptide by alterations in primary sequence. These include genetic variants, both natural and induced (for example, resulting from random mutagenesis by irradiation or exposure to ethanemethylsulfate or by site-specific mutagenesis as described in Sambrook *et al.*, *supra*; or Ausubel *et al.*, *supra*). Also included are cyclized peptides, molecules, and analogs which contain residues other than L-amino

25 acids, *e.g.*, D-amino acids or non-naturally-occurring or synthetic amino acids, *e.g.*, B or Y amino acids. In addition to full-length polypeptides, the invention also includes SAP family member polypeptide fragments. As used herein, the term "fragment," means at least 20 contiguous amino acids, preferably at least 30 contiguous amino acids, more preferably at least 50 contiguous amino acids, and most preferably at least

-77-

60 to 80 or more contiguous amino acids. Polypeptide fragments of SAP family members can be generated by methods known to those skilled in the art or may result from normal protein processing (*e.g.*, removal of amino acids from the nascent polypeptide that are not required for biological activity or removal of amino acids by  
5 alternative mRNA splicing or alternative protein processing events).

Preferable fragments or analogs according to the invention are those which facilitate specific detection of a SAP family member nucleic acid or amino acid sequence in a sample to be diagnosed. Particularly useful SAP family member polypeptide fragments for this purpose include, without limitation, the amino acid  
10 fragments corresponding to the SH2 domain.

All publications and patent applications mentioned in this specification are herein incorporated by reference to the same extent as if each independent publication or patent application was specifically and individually indicated to be incorporated by reference.

15 While the invention has been described in connection with specific embodiments thereof, it will be understood that it is capable of further modifications and this application is intended to cover any variations, uses, or adaptations of the invention following, in general, the principles of the invention and including such departures from the present disclosure come within known or customary practice  
20 within the art to which the invention pertains and may be applied to the essential features hereinbefore set forth, and follows in the scope of the appended claims.

Other embodiments are within the claims.

What is claimed is:

-78-

Claims

1. A substantially pure nucleic acid encoding a SAP polypeptide.
2. The nucleic acid of claim 1, wherein said SAP polypeptide is a fragment of the full length naturally-occurring SAP polypeptide.
- 5           3. The nucleic acid of claim 1, wherein said nucleic acid is operably linked to a second nucleic acid.
4. The nucleic acid of claim 3, wherein said second nucleic acid is a coding sequence and said nucleic acid operably linked to said second nucleic acid produces a fusion protein comprising said SAP polypeptide.
- 10           5. The nucleic acid of claim 3, wherein said second nucleic acid is a gene promoter.
6. The nucleic acid of claim 1, wherein said nucleic acid comprises a mutation, said mutation resulting in an amino acid alteration in said SAP polypeptide.
7. The nucleic acid of claim 1, wherein said nucleic acid comprises a nucleic  
15   acid sequence that is substantially identical to SEQ ID NO: 3 or is substantially identical to SEQ ID NO: 5.
8. The nucleic acid of claim 1, wherein said nucleic acid comprises a nucleic acid sequence encoding a naturally-occurring SH2 domain.
9. The nucleic acid of claim 1, wherein said polypeptide has SAP biological  
20   activity.



-79-

10. The nucleic acid of claim 1, wherein said polypeptide modulates SH2 domain-containing protein-mediated signal transduction.

11. A substantially pure SAP polypeptide.

12. The polypeptide of claim 11, wherein said polypeptide is a fragment of  
5 the full length naturally-occurring SAP polypeptide.

13. The polypeptide of claim 11, wherein said polypeptide is part of a fusion protein.

14. The polypeptide of claim 11, wherein said polypeptide comprises an amino acid sequence that has a mutation as compared to the naturally-occurring  
10 amino acid sequence of said polypeptide.

15. The polypeptide of claim 11, wherein said polypeptide comprises an amino acid sequence that is substantially identical to SEQ ID NO: 4 or is substantially identical to SEQ ID NO: 6.

16. The polypeptide of claim 11, wherein said polypeptide comprises a  
15 naturally-occurring SH2 domain.

17. The polypeptide of claim 11, wherein said polypeptide has SAP biological activity.

18. The polypeptide of claim 11, wherein said polypeptide modulates SH2 domain-containing protein-mediated signal transduction.

20 19. A method of treating a disease involving aberrant SH2 domain-containing

-80-

protein-mediated signal transduction in a patient, said method comprising administering to said patient a SAP family member polypeptide or a fragment, mutant, or fusion thereof.

20. The method of claim 19, wherein said polypeptide comprises an amino acid sequence substantially identical to SEQ ID NO: 4 or is substantially identical to SEQ ID NO: 6.

21. The method of claim 19, wherein said polypeptide is an EAT-2 polypeptide.

22. A method of identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction, said method comprising:

- (a) providing a cell comprising a SAP family member polypeptide-encoding gene;
- (b) contacting said cell with a candidate compound; and
- (c) monitoring expression of said SAP family member polypeptide-encoding gene, an alteration in the level of said expression of said gene in response to said candidate compound indicating the presence of a compound that modulates SH2 domain-containing protein-mediated signal transduction.

23. A method of identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction, said method comprising:

- (a) providing a cell comprising a reporter gene operably linked to a promoter from a SAP family member polypeptide-encoding gene;
- (b) contacting said cell with a candidate compound; and
- (c) measuring expression of said reporter gene, an alteration in the level of said expression of said reporter gene in response to said candidate compound indicating the presence of a compound that modulates SH2 domain-containing

-81-

protein-mediated signal transduction.

24. The method of claim 22 or 23, wherein said SAP family member polypeptide-encoding gene encodes a SAP polypeptide or encodes an EAT-2 polypeptide.

- 5           25. A method of identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction, said method comprising:
- (a) providing a cell having:
- (i) a reporter gene operably linked to a DNA-binding-protein recognition site;
- 10               (ii) a first fusion gene capable of expressing a first fusion protein, said first fusion protein comprising a SAP family member polypeptide covalently bonded to a binding moiety, said binding moiety capable of specifically binding to said DNA-binding-protein recognition site; and
- (iii) a second fusion gene capable of expressing a second fusion
- 15           protein, said second fusion protein comprising a SLAM polypeptide covalently bonded to a gene activating moiety;
- (b) exposing said cell to a candidate compound; and
- (c) measuring reporter gene expression in said cell, an alteration in the level of said expression of said reporter gene in response to said candidate compound
- 20           indicating the presence of a compound that modulates SH2 domain-containing protein-mediated signal transduction.

26. A method of identifying a compound that modulates SH2 domain-containing protein-mediated signal transduction, said method comprising:

- (a) providing a cell having:
- 25               (i) a reporter gene operably linked to a DNA-binding-protein recognition site;

-82-

(ii) a first fusion gene capable of expressing a first fusion protein, said first fusion protein comprising a SLAM polypeptide covalently bonded to a binding moiety, said binding moiety capable of specifically binding to said DNA-binding-protein recognition site; and

5 (iii) a second fusion gene capable of expressing a second fusion protein, said second fusion protein comprising a SAP family member polypeptide covalently bonded to a gene activating moiety;

(b) exposing said cell to said compound; and

(c) measuring reporter gene expression in said cell, an alteration in the level  
10 of said expression of said reporter gene in response to said candidate compound indicating the presence of a compound that modulates SH2 domain-containing protein-mediated signal transduction.

27. A method of identifying a compound as an SAP family member protein mimetic, said method comprising the steps of:

15 (a) providing a SLAM polypeptide, or a fragment or fusion thereof, said SLAM polypeptide bearing non-phosphorylated tyrosine residues;

(b) contacting said SLAM polypeptide with a SAP family member polypeptide;

(c) contacting said SLAM polypeptide and said SAP family member  
20 polypeptide with a candidate compound; and

(d) measuring level of interaction of said SLAM polypeptide with said SAP family member polypeptide, a decrease in said level in response to said compound relative to a level not contacted with said compound indicating said compound is a SAP family member protein mimetic.

25 28. The method of claim 25, 26, or 27, wherein said SAP family member polypeptide is a SAP protein or an EAT-2 protein.

-83-

29. A method of identifying a compound as an SAP family member protein mimetic, said method comprising the steps of:

(a) providing a SLAM polypeptide, or a fragment or fusion thereof, said SLAM polypeptide bearing non-phosphorylated tyrosine residues; and

5 (b) contacting said SLAM polypeptide with a candidate compound, wherein said candidate compound binding to said SLAM polypeptide indicates said candidate compound is a SAP family member protein mimetic.

30. The method of claim 27 or 29, wherein said SLAM polypeptide is bound to a solid state substrate.

10 31. A method of identifying a polypeptide that modulates SH2 domain-containing protein-mediated signal transduction, said method comprising:

(a) providing a cell having:

(i) a reporter gene operably linked to a DNA-binding-protein recognition site;

15 (ii) a first fusion gene capable of expressing a first fusion protein, said first fusion protein comprising a SAP family member polypeptide covalently bonded to a binding moiety, said binding moiety capable of specifically binding to said DNA-binding-protein recognition site; and

(iii) a second fusion gene capable of expressing a second fusion  
20 protein, said second fusion protein selected from a library, said library comprising a polypeptide covalently bonded to a gene activating moiety, said polypeptide encoded by a cDNA of said library; and

(b) measuring reporter gene expression in said cell, an increase in said  
25 reporter gene expression identifying the presence of a polypeptide that modulates SH2 domain-containing protein-mediated signal transduction.

32. A method of modulating SH2 domain-containing protein-mediated signal

-84-

transduction in a mammal, said method comprising providing a transgene encoding a SAP family member polypeptide or fragment thereof to a cell of said mammal, said transgene being positioned for expression in said cell.

33. A method of modulating SH2 domain-containing protein-mediated signal  
5 transduction in a mammal, said method comprising administering to a cell of said mammal a compound which modulates the biological activity of a SAP family member polypeptide.

34. A method of increasing antigen-specific T cell activation in a mammal,  
said method comprising providing a transgene encoding a SAP family member  
10 polypeptide, said transgene being positioned for expression in said cell.

35. A method of increasing antigen-specific T cell activation in a mammal,  
said method comprising administering to a cell of said mammal a compound which  
increases the biological activity of a SAP family member polypeptide.

36. The method of claim 34 or 35, wherein said antigen-specific T cell  
15 activation is mediated by Th1 helper T cells.

37. The method of claim 31, 32, 33, 34, or 35, wherein said SAP family  
member polypeptide is a SAP protein or an EAT-2 protein.

38. A transgenic mammal, said mammal having a knockout mutation in an  
endogenous SAP family member protein-encoding nucleic acid sequence.

20 39. A transgenic mammal, said mammal having an exogenous SAP family  
member protein-encoding nucleic acid sequence operably linked to a promoter.

-85-

40. The transgenic mammal of claim 38 or 39, wherein said SAP family member protein is a SAP protein or an EAT-2 protein.

41. The transgenic mammal of claim 38 or 39, wherein said animal has aberrant SH2 domain-containing protein mediated signal transduction.

```
1 / 1 31 / 11
cag ttg aga aga aga ggt aaa acg aac cat tac cag aca aca gtg gaa aaa aaa agc ctt
gln leu arg arg arg gly lys thr asn his tyr gln thr thr val glu lys lys ser leu
61 / 21 91 / 31
acg atc tat gcc caa gtc cag aaa cca ggt cct ctt cag aag aaa ctt gac tcc ttc cca
thr ile tyr ala gln val gln lys pro gly pro leu gln lys lys leu asp ser phe pro
121 / 41 151 / 51
gct cag gac cct tgc acc acc ata tat gtt gct gcc aca gag cct gtc cca gag tct gtc
ala gln asp pro cys thr thr ile tyr val ala ala thr glu pro val pro glu ser val
181 / 61 211 / 71
cag gaa aca aat tcc atc aca gtc tat gct agt gtg aca ctt cca gag agc
gln glu thr asn ser ile thr val tyr ala ser val thr leu pro glu ser
```

Figure 1



## HUMAN SAP

## DNA

GGTTGACTTGTGCCTGGCTGCAGTAGCAGCGGCATCTCCCTTGCACAGTTCTCCTCCTCGG  
CCTGCCCAAGAGTCCACCAGGCCATGGACGCAGTGGCTGTGTATCATGGCAAATC  
AGCAGGGAAACCGGCGAGAAGCTCCTGCTTGCCACTGGGCTGGATGGCAGCTAT  
TTGCTGAGGGACAGCGAGAGCGTGCCAGGCGTGTACTGCCTATGTGTGCTGTAT  
CACGGTTACATTTATACATACCGAGTGTGCCAGACAGAAACAGGTTCTTGGAGT  
GCTGAGACAGCACCTGGGGTACATAAAAGATATTTCCGGAAAAATAAAAAATCTC  
ATTTCAGCATTTTCAGAAGCCAGATCAAGGCATTGTAATACCTCTGCAGTATCCA  
GTTGAGAAGAAGTCCTCAGCTAGAAGTACACAAGGTACTACAGGGATAAGAGAA  
GATCCTGATGTCTGCCTGAAAGCCCCATGAAGAAAAATAAACACCTTGTACTTTAT  
TTTCTATAATTTAAATATATGCTAAGTCTTATATATTGTAGATAATACAGTTCGGTGAGCT  
ACAAATGCATTTCTAAAGCCATTGTAGTCCGTGAATGGAAGCATCTAGCATGTCGTCAAAG  
CTGAAATGGACTTTTGTACATAGTGAGGAGCTTTGAAACGAGGATTGGGAAAAAGTAATTC  
CGTAGGTTATTTTCAGTTATTATATTTACAAATGGGAAACAAAAGGTAATGAATACTTTAT  
AAAGGATTAATGTCAATTCTTGCCAAATATAAATAAAAAATAATCCTCAGTTTTTGTGAAAA  
GCTCCATTTTTAGTGAAATATTATTTATAGCTACTAATTTTAAATGTCTTGCTTGATTG  
TATGGTGGGAAGTTGGCTGGTGTCCCTTGTCTTTGCCAAGTCTCCACTAGCTATGGTGTCT  
ATAGGCTCTTTTGGGATTTTTGAAGCTGTATACTGTGTGCTAAACAAGCACTAAACAAAG  
AGTGAAGGATTTTATGTTTAAATCTGAAAGCAACCTTCTTGCCTAGTGTCTGATATTGGAC  
AGTAAATCCACAGACCAACCTGGAGTTGAAAATCTTATAATTTTAAATATGCTCTAAACA  
TGTTTATCGTATTTGATGCTACAGGATTTGAAATTGTATTACAAATCCAATGAATAGAGTT  
TTTCTTTTCATTTACCTCTGCCCCAGTTGTTTCTACTACATGGAAGACCTCATTTTGAAGG  
GAAATTTTCAGCAGCTGCAGCTCATGAGTAAGTGAATTTGTAACAAGCCTCCTTTTAAAGTAA  
CCCTACAAAACCACTGGAAAGTTTATGGTTGTATTATTTTTTAAAAAAATTCGAAGTGATT  
GAAACCTACACGAGATACAGAATTTTATGCGGCATTTTCTTCTCACATTTATATTTTTGTG  
ATTTTGTGATTGATTATATGTCACCTTTTGCTACAGGGCTCACAGAATTCATTCACCTCAACAA  
ACATAATAGGGCGCTGAGGGCATAGAAGTAAAAACACCTGGTCCCTGCTCTCAGTTCAGTG  
TCTTGTTGGACGAGAAAAACAATAACGATAAAAGACAGTGAAAGAAATAACGATAAAAGAC  
AGTGAAAGAAAAATAACAATAAAAGACAAGGAAAAATAACAATGAAAGTTGATAAGTACAT  
GATAAGCGAGGTTCCCCGTGTGTAGGTAGATCTGGTCTTTAGAGCCAGATAGATAGGTCAG  
TGCAAATACTCTGGTCCATGGGCCATATGAAAAGGCTAAGCTTCACTGTAAAATAATAACT  
GGGAATTCTGGATTGTGTATGGGTGTTGGTGAACCTGGGTTTTAATTAGTGAACCTGCTGAGA  
GACAGAGCTATTCTCCATGTACTGGCAAGACCTGATTTCTGAGCATTAAATATGGATGCCG  
TGGGAGTACAAAAGTGGAGTGTGGCCTGAGTAATGCATTATGGGTGGTTTACCATTTCCTTG  
AGGTAAAAGCATCACATGAACCTGTAAAGGAATTTAAATACTCTTCATAATAAGTTG  
CATAGGTTTAAATAATTTTAAATATATGGCTTGAGTTTAAATTGTAATAGGCGTAACTAAT  
TTTAACTCTATAATGTGTTCATCTGGAATAATCCTAAACATATGAATTATGTTTGCATGT  
TCACTTCCAAGAGCCTTTTTTTGAAAAAAGCTTTTTTTGAATCATCAAGTCTTTCACATT  
TAAATAAAGTGTTTGAAAGC

Figure 2A

## PROTEIN

MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSSEVPGVYCLCVLYHGYIYTYRVSQTET  
GSWSAETAPGVHKRYFRKIKNLI SAFQKPDQGIVIPLOYPEKKSSARSTQGTGTGIREDPD  
VCLKAP

Figure 2B

**SAP Sequences:****mouse SAP cDNA:**

ATGGATGCAGTGACTGTGTACCACGGCAAAATCAGCAGGGAGACCGGGGAGAAGCTCTTACTCGCTACCGGGCTGGAT  
GGAAGCTATCTGCTGCCGAGACAGCGAGAGTGTCCCTGGCGTGTA CTGCCTGTGTGTTTTGTATCAAGGTTACATCTAC  
ACATATCSAGTGTCCCAGACAGAAACAGGTTCTTGGAGTGCCGAGACAGCACCTGGAGTACATAAAAGATTTTCCGG  
AAAGTAAAAATCTCATCTCAGCGTTTCAGAAGCCGGATCAAGGCATCGTGACGCCCTCTGCAGTATCCAGTTGAAAAG  
TCCTCTGGCAGGGGCCACAAGCTCCACAGGGAGAAGAGATTCTGATATCTGCCTGAATGCACCATGA

Figure 2C

**mouse SAP protein:**

MDAVTVYHGKISRETGERLLLATGLDGSYLLRDSESVPGVYCLCVLYQGYIYTYRVSQTETGSWSAETAPGVHRRFFR  
KVKNLISAFQKPDGGIVTPLQYPVEKSSGRGPGAPTGRDSDICLNAP

Figure 2D

**Mutants****hSAPΔ55 cDNA:**

ATGGACGCAGTGGCTGTGTATCATGGCAAAATCAGCAGGGAAACCGGCGAGAAGCTCCTG  
CTTGCCACTGGGCTGGATGGCAGCTATTTGCTGAGGGACAGCGAGAGCGTGCCAGGCGTG  
TACTGCCTATGTGTGCTGTATCACGGTTACATTTATACATACCGAGTGTCCCAGACAGAA  
ACAGGTTCTTGGAGTGCTGAGCATTTCAGAAGCCAGATCAAGGCATTGTAATACCTCTGC  
AGTATCCAGTTGAGAAGAAGTCCTCAGCTAGAAGTACACAAGGTACTACAGGGATAAGAG  
AAGATCCTGATGTCTGCCTGAAAGCCCCATGA

**hSAPΔ55 protein:**

MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSESVPGVYCLCVLYHGYIYTYRVSQTE  
TGSWSAEHFRSQIKAL

**A1-1 cDNA:**

ATGGACGCAGTGGCTGTGTATCATGGCAAAATCAGCAGGGAAACCGGCGAGAAGCTCCTG  
CTTGCCACTGGGCTGGATGGCAGCTATTTGCTGAGGGACAGCGAGAGCGTGCCAGGCGTG  
TACTGCCTATGTGTGCTACAGCACCTGGGGTACATAAAAGATATTTCCGAAAATAAAAA  
ATCTCATTTTCAGCATTTTCAGAAGCCAGATCAAGGCATTGTAATACCTCTGCAGTATCCAG  
TTGAGAAGAAGTCCTCAGCTAGAAGTACACAAGGTACTACAGGGATAAGAGAAGATCCTG  
ATGTCTGCCTGAAAGCCCCATGA

**A1-1 protein:**

MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSESVPGVYCLCVLQHLGYIKDISGK

**A1-2 cDNA:**

ATGGACGCAGTGGCTGTGTATCATGGCAAAATCAGCAGGGAAACCGGCGAGAAGCTCCTG  
CTTGCCACTGGGCTGGATGGCAGCTATTTGCTGAGGGACAGCGAGAGCGTGCCAGGCGTG  
TACTGCCTATGTGTGCTCATTTTCAGAAGCCAGATCAAGGCATTGTAATACCTCTGCAGTA  
TCCAGTTGAGAAGAAGTCCTCAGCTAGAAGTACACAAGGTACTACAGGGATAAGAGAAGA  
TCCTGATGTCTGCCTGAAAGCCCCATGA

**A1-2 protein:**

MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSESVPGVYCLCVLISEARSRHCNTSAVSS

**Exon2 genomic mutant:**

ggaaactgtggttgggcagatacaatatggagacaagtttagattcactatgctttacttc  
ctatgaatgcaatgacaccatatacgtgtgtcctagtatatgtgacatttatatgtaata  
ttaagctcaaatttaaagtatccattgttcttttggaaatctttcagtaatggaagtttat  
tctttcagGTATCACGGTTACATTTATACATACCGAGTGTCCCAGACAGAAACAGGTT  
CTTGAGTGTGAGgtatagttgtattttatTTTTgttctgtgggggtgtcaaggaggtatt  
tgaaatttaggctggttttataaaagagcaaattatacattattaagattcataaggtt  
taaattctctaaagctccaatccaaaattgttcatggatcattaagaaagcttttagagaat  
tttggtcccagtcctggttagcc

**Figure 3**

*The amino acid sequence of human SAP with the consensus motifs of SH2 domains in bold*

10	20	30	40	49
MDAVAVYHGKISRETGEKLL	ATGLDGSYLLRDS	ESVPGVYCLCVLYHG		
60	70	80	90	100
YIYTYRVSQTETG	SWSAETAPGVH	KRYFRKIKNLISAFQ	KPDQGIVPLQY	
110	120	128		
PVEKKSSARSTQ	GTTGIREDPD	VCLKAP		

Figure 4A

*Consensus motifs found in SH2 domains as defined by a program termed BLOCKS, i.e. BLOCK PR00401: SH2 DOMAIN*

Internet Address: [www.blocks.fhcrc.org](http://www.blocks.fhcrc.org)  
 Reference: S Henikoff and JG Henikoff "Protein family classification based on searching a database of blocks" *Genomics* 19: 97 (1994)

BLOCK PR00401A	VYHGKISRETGEKLL	SAPh
BLOCK PR00401B	DGSYLLRDS	SAPh
BLOCK PR00401C	PGVYCLCVLYHG	SAPh
BLOCK PR00401D	YIYTYRVSQTE	SAPh
BLOCK PR00401E	YFRKIKNLISAFQKP	SAPh

Figure 4B

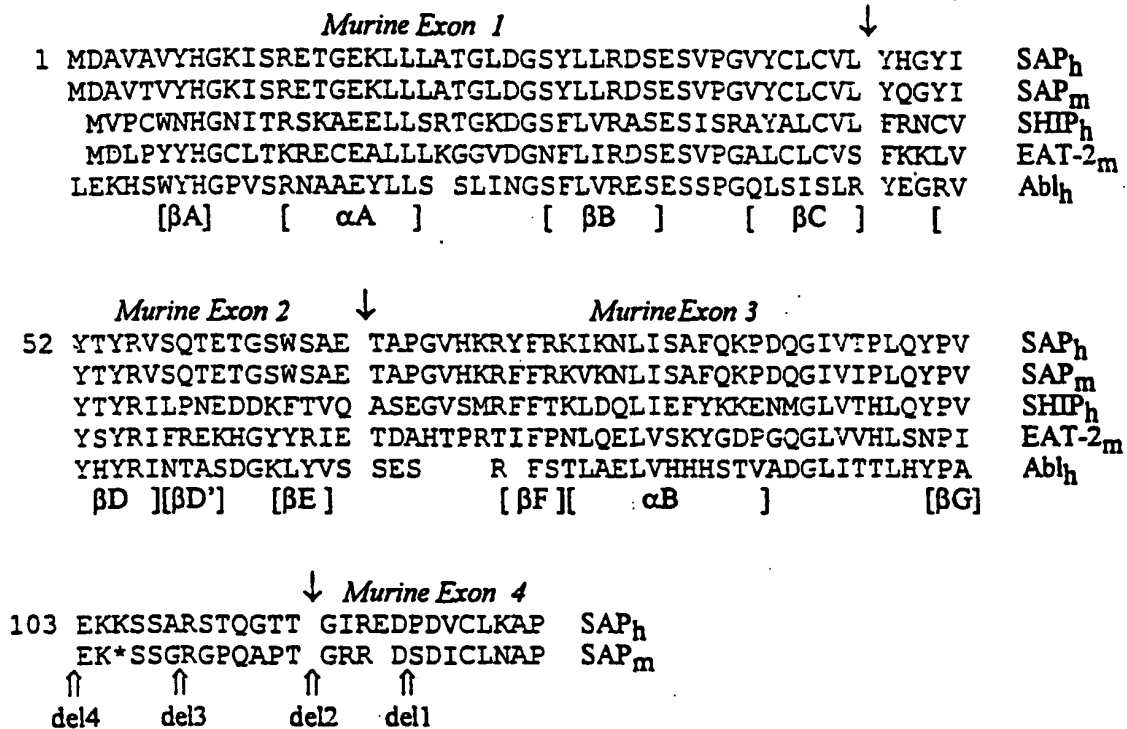


Figure 5

Figure 6A

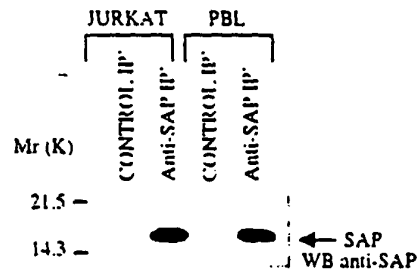


Figure 6C

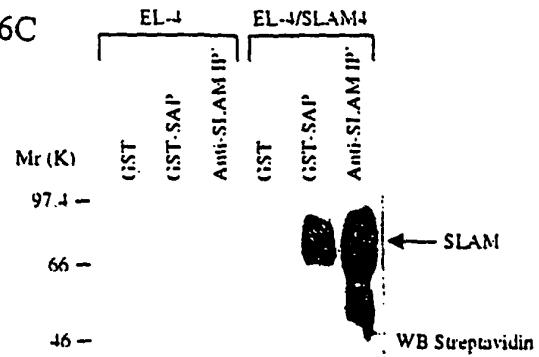


Figure 6B

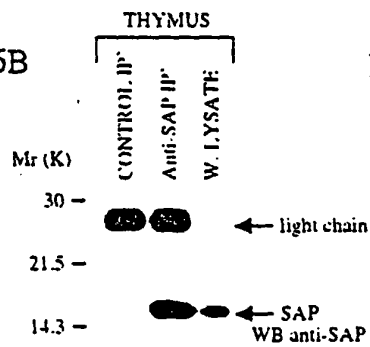
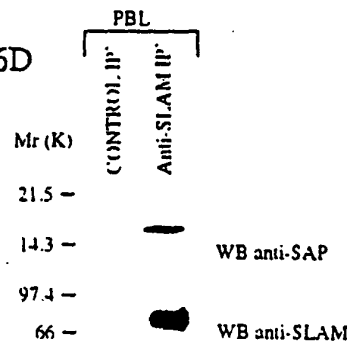


Figure 6D



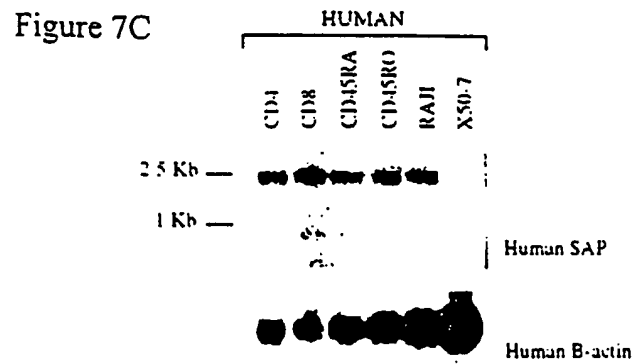
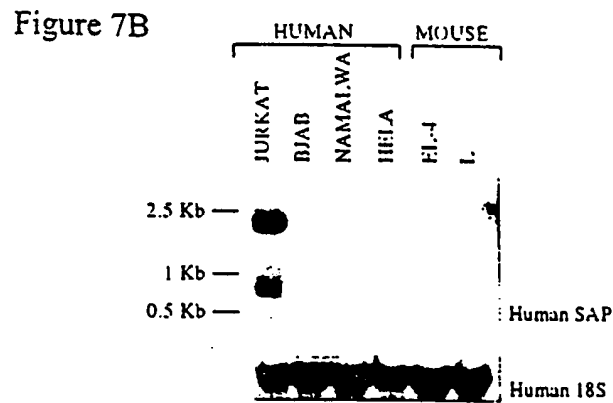
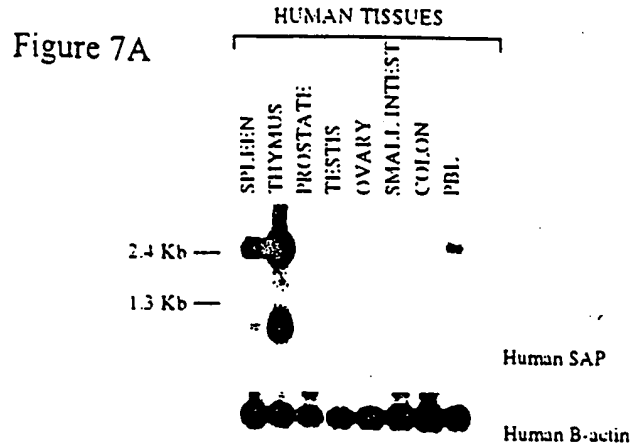


Figure 8A

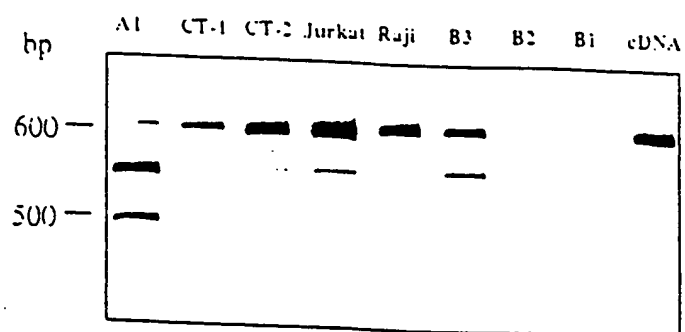


Figure 8B

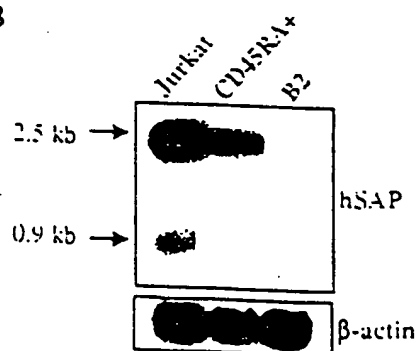




Figure 9A

```

87 ATGGACGCASTGGCTGTGTATCATGGCAAAATCAGCAGGGAAACCGGCCGAGAGCTCCTGCTT hSAP
150 GCCACTGGGCTGGATGGCAGCTATTTGCTGAGGGACAGCGAGACCGTGCCAGCCGTGTACTGC hSAP
213 CTA TGT GTG CTG TAT CAC GGT TAC ATT TAT ACA TAC CGA GTG TCC CAG hSAP
    CTA TGT GTG CTG TAT CAC GGT TAC ATT TAT ACA TAC CGA GTG TCC CAG hSAPΔ55
    CTA TGT GTG CT- --- --- --- --- --- --- --- --- --- A1-1
    CTA TGT GTG CT- --- --- --- --- --- --- --- --- --- A1-2
261 ACA GAA ACA GGT TCT TGG AGT GCT GAG ACA GCA CCT GGG GTA CAT AAA hSAP
    ACA GAA ACA GGT TCT TGG AGT GCT GAG --- --- --- --- --- hSAPΔ55
    --- --- --- --- --- --- --- --- --- ACA GCA CCT GGG GTA CAT AAA A1-1
    --- --- --- --- --- --- --- --- --- --- --- --- --- A1-2
309 AGA TAT TTC CGG AAA ATA AAA AAT CTC ATT TCA GCA TTT CAG AAG CCA hSAP
    --- --- --- --- --- --- --- --- --- --- --- -CA TTT CAG AAG CCA hSAPΔ55
    AGA TAT TTC CGG AAA ATA AAA AAT CTC ATT TCA GCA TTT CAG AAG CCA A1-1
    --- --- --- --- --- --- --- --- --- --- --- -CA TTT CAG AAG CCA A1-2
357 GATCAAGGCATTGTAATACCTCTGCAGTATCCAGTTGAGAAGAAGTCCTCAGCTAGAAGTACA hSAP
420 CAAGGTACTACAGGGATAAGAGAAGATCCTGATGTCTGCCTGAPAGCCCCATGA hSAP

```

Figure 9B

```

EXON 1
1 MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSESVPGVYCLCVL YHGVI hSAP
  MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSESVPGVYCLCVL YHGVI hSAPΔ55
  MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSESVPGVYCLCVL QHLGY A1-1
  MDAVAVYHGKISRETGEKLLLATGLDGSYLLRDSESVPGVYCLCVL TSEAR A1-2

EXON 2 EXON 3
YTYRVSQTETGSWSAE TAPGVHKRYFRKIKNLISAFQKPDQGIPTPLQYPV hSAP
YTYRVSQTETGSWSAE HFPSSQIKAL 76 hSAPΔ55
IKDISGK 58 A1-1
SRHCNTSAVSS 62 A1-2

EXON 4
EKKSSARSTQGTG GIREDPDVCLKAP 128 hSAP

```

Figure 10A

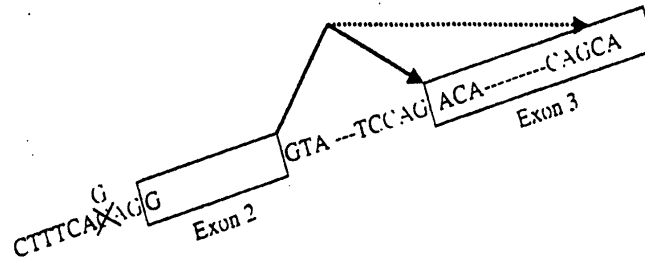


Figure 10B

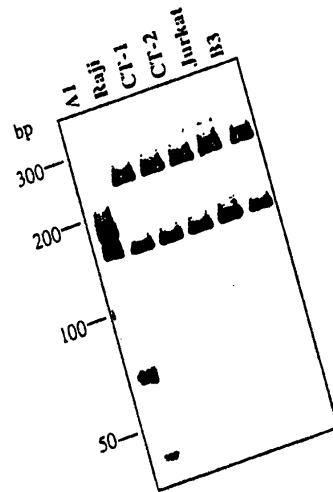
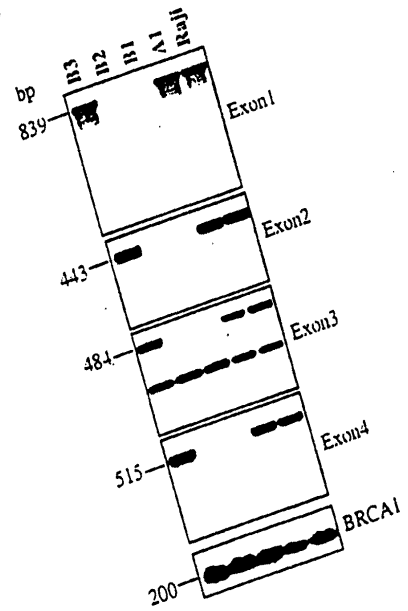
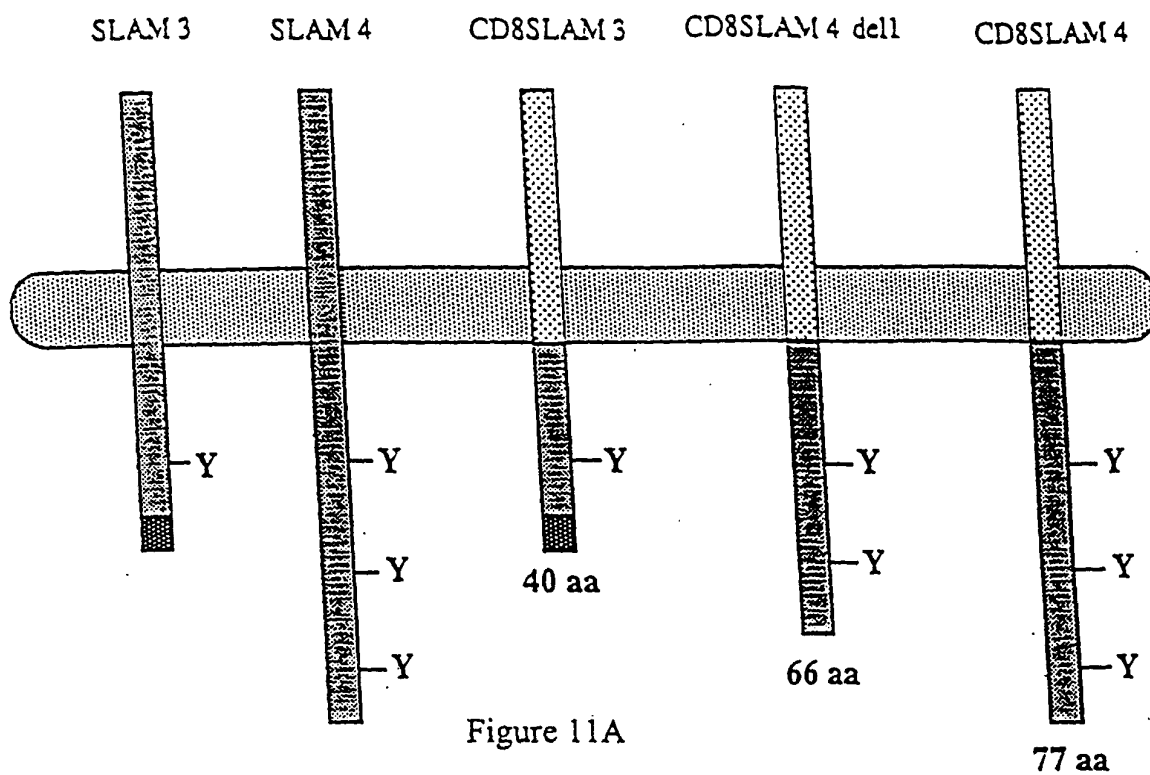


Figure 10C



## CD8SLAM PROTEINS



SLAM 3	QLRRRGKTNHYQTTVEKKSLTIYAQVQKPGDTHHQTSDLF
SLAM 4 del 1	QLRRRGKTNHYQTTVEKKSLTIYAQVQKPGPLQKKLDSFPA QDPCTTIYVAATEPVPESVQETNSI
SLAM 4	QLRRRGKTNHYQTTVEKKSLTIYAQVQKPGPLQKKLDSFPA QDPCTTIYVAATEPVPESVQETNSITVYASVTLPE

Figure 11B

Figure 12A

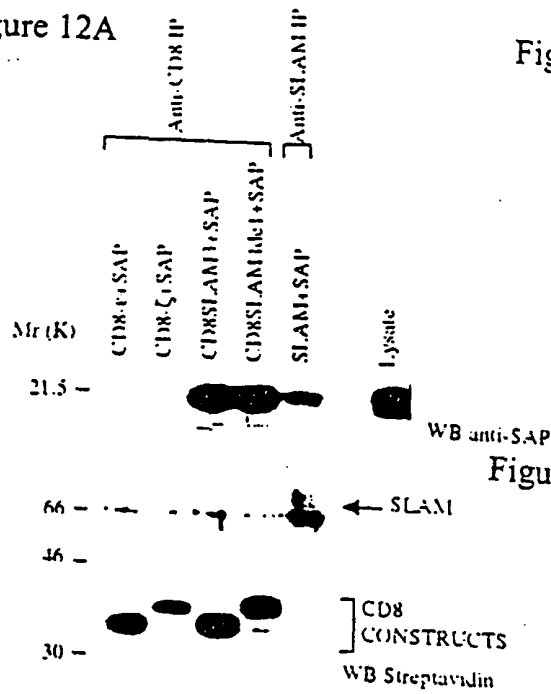


Figure 12C

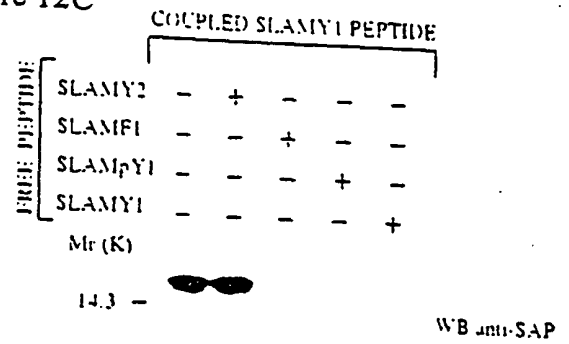


Figure 12D

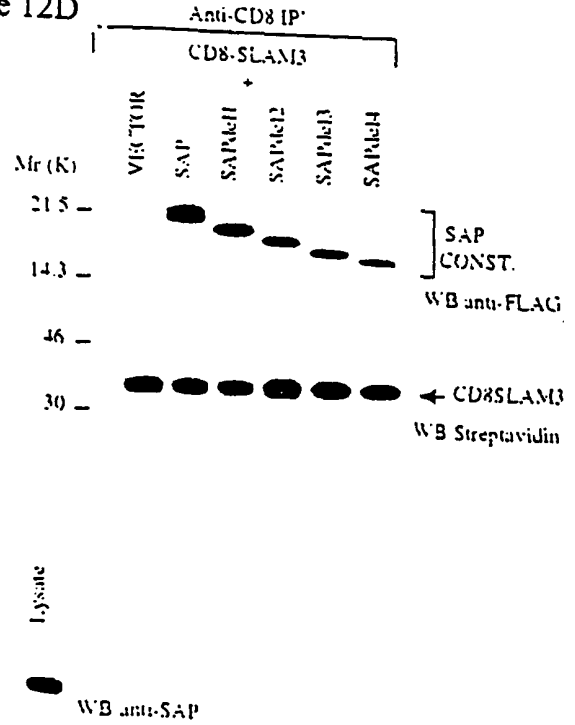


Figure 12B

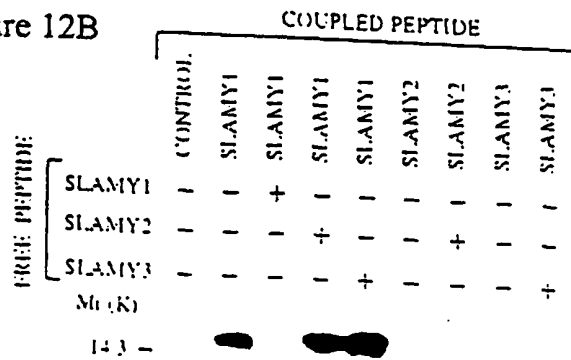


Figure 13A

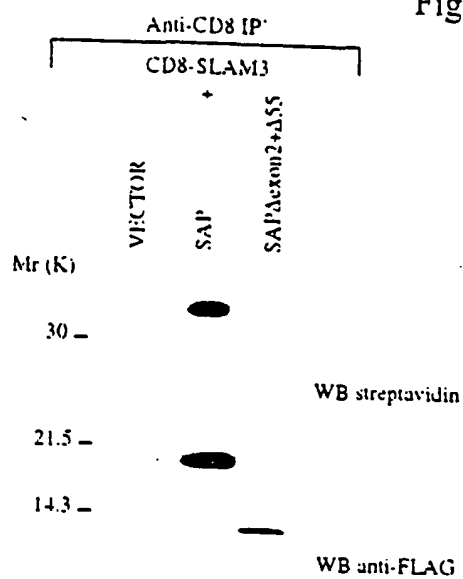


Figure 13C

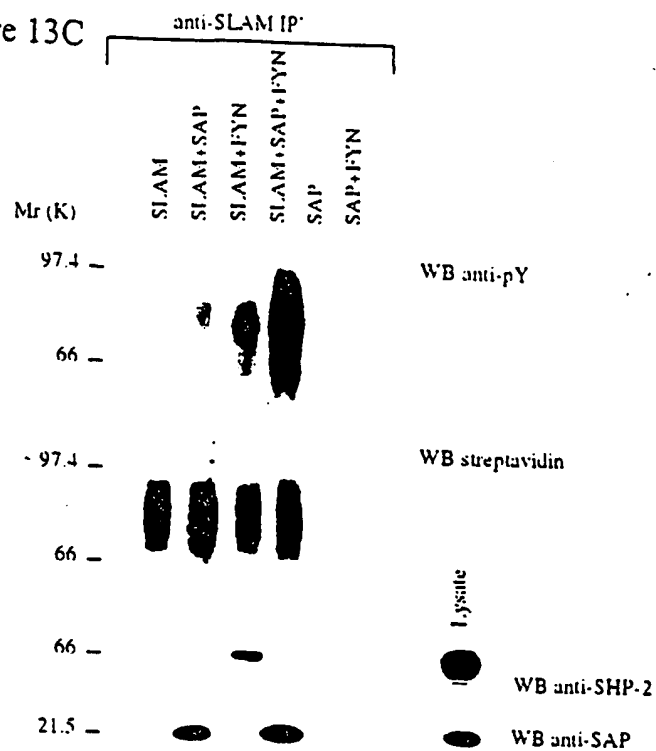


Figure 13B

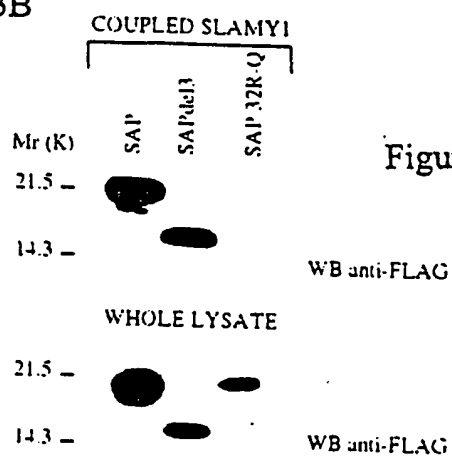
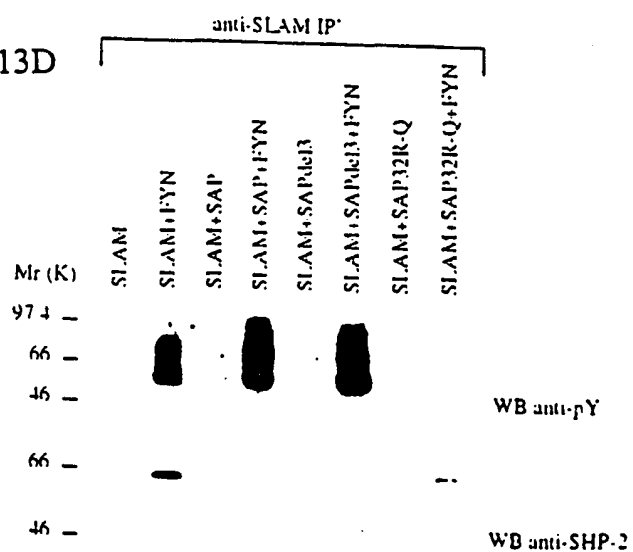


Figure 13D



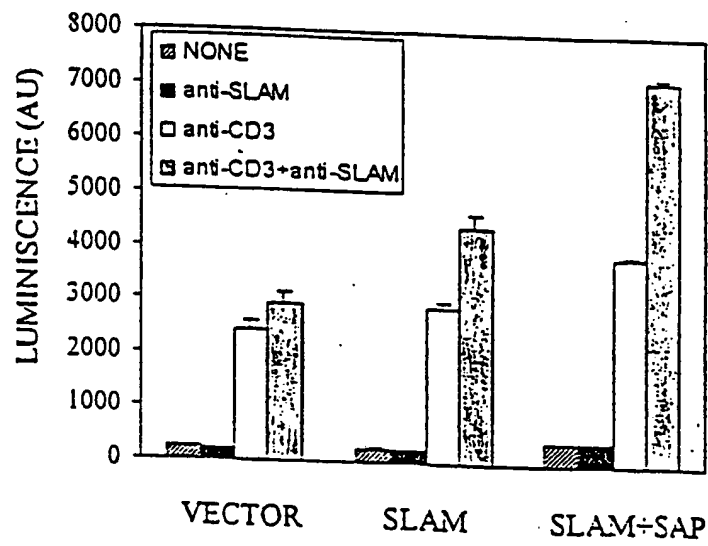


Figure 14



CAGCAAAGTTTCCTTTTACCATCCTTCTTCTGGGCCCATCAAAGASTTTGCTAGGCATGG  
ATGCAGTGACTGTGTACCACGGCAAATCAGCAGGGAGACCGGGGAGAAGCTCTTACTCGC  
TACCGGGCTGGATGGAAGCTATCTGCTGCGAGACAGCGAGAGTGTCCCTGGCGTGTACTGC  
CTGTGTGTTTTgtgagtatatctcgtggtagtggcaggttggttagcatgggtagcctgccc  
tttcaggggagggctgggaggagtggcaggggatgacagcttttggcagaggtacctaggg  
aatacaggcagcaggaactgagtactcttccagctctgacgtatggtcccagggctcagcc  
cagggatgtgggtgggcagtacttgtacgtacgtccctcactgcggacttttatcaactgcat  
ttcaccagtgcgtttgtggcaacttcactgaatacccttcagaatgggaaaacatggccta  
gaacttacctttactttgagaaagacccatactctggacttcggctctgattccaagtaga  
caccatgatggtggagccccgggaagttggactatttgtattctgggcatgaagtaaggc  
aaaggaaggtctttctctatcattctctacaaatgtgactgaaaatggtagccctaaaagaa  
aagcatctagcaaatgcttgaagagtgggttagccaggaaatggaatccattgctttcagaa  
tgcagcccaggggaaggcagaggttagacagatgtgtgtgagtttcaggtgactaacctgggt  
ttacatggcccggtcatccaactaggctacatagataaaaaaccaag



## EXON2 / INTRON2

GAAACCCCTTGGAGACCAAACAGGTTCTTGGAGTGCCGAggtacagctcttttatttggtt  
ttacttctctgggtataagaaaagtgccttgaaatctaggttggtttaaaaaagagcagtca  
tactatacactaccaaaatatttcaatcactaatgctcaattctatagtcattgatggttc  
atrtagaagccttttagaaaatttggttccagacttatttagccaccattatgtgaggtag  
gatcattttgggaaaatgaaaaaatttgaaggaccatgtgtaacttatagtgctctttacaa  
acttttagaaagcatataacaataataacaacaacaacaacaacaacaacaataa  
taatacaaacctaaaaaactaaaaaaaaaaaaaaaaagggaaccatatactaaatgatactg  
atgcacagatttggttctgttggtgataaagcaaacctgggatttctatttctactagtact  
agtaccccttttggaactcctaacaggggattatactacagcgtgaaatcctgggcatgtca  
aatgcattcagactccatgtgtagttcttctctatagtcacttttgatggaagcaatttgta  
aaatgggtctacaaacatttatcaagaggaccatttcaggaggtataaggcaatgcaac  
taaagctcttagctggagagctctgacatacagtgagcatccaaaaatgtgaactactgaaa  
acttctcagcagtatgattgtgtgtcatctgggtgtaaatttaccaggcacaagaatttgaa  
cagaggcattttcaaacatccaggcgattgctgctgctctgtgtgcataggcaagggtttc  
aaaggagccataaaagagcctctggatagtcggaaaagagaagagggaatttgtagttt  
tatttaataaaaaatcttctttgagaatttcaaatatatacacaaatgaattcctatctgc  
tattgtagtgaatttaaaaggatgtttatagaatgatgttctcctccatcttgggactccc  
taggagtcgaagaagaagaaggaggaggtgagtggtatacttctgggtaccaatcttg  
gcttctactgggttagatcccttttagtttctcttatgaactagggctttgaataacattaat  
tcagaatgaagggttcttatggccttgaaatatttgtacaccactgtgacagtcagaaaat  
tattgaactatttctagagagcaggtgggtatagagcaatgcatttatgaatcaattcatga  
ttctacagctgtggttttcaaacctttaggccttttagttggttctgaccttatactagttta  
tcttgatagtggaatgcttcacaatgatcttttgaaagcatagatgtcaatatcaatttct  
tattagtaactaaataaatctgtgcatttaaaagaataataaagtcctagacaaaagtttaa  
gaatgtcttacaattttgtcagtaaaaagcaaggtataaatttcaagttatttgaaattatca  
aaacaaattagaatatattcaagtatactgttatcttatttaattgggttaattattttta  
agttttatttttaactttcagggttcgtaaaaaatagatgaaggatgaaaccagggtctca  
tgcattttagacaattgtttataccattgaataatattaccaataatttctctttttattg  
ccatatattcgctgtacatgttaatgcacttcataagacaagtttataatatcttttgatt  
atgttctctcataaaatactatatccctttgtggccatgttggttattttgtgtcaggatac  
tatccaagggaataatcaataccatttctaataaaaaactatacttcttaacaaatctatca  
ctttaaataaataagcaatttgacttatcatgaataataattatgtatagctgatacatat  
gaatttggtgcatgggttgctatatatttaactcttttccacccaagaaatagccttcagatg  
agcaatagaaaatttaattggagagcctttgatttttttttttacttaaaaaatagttct  
ttgagaattttgtacaatatgttttgatcatatttacctttgtcttctaactccttcagat  
ccacccttccctctctattcaccgccttttatactcctgtttatccctaagctgtgcaat  
gcagtttggtgctacccaaatattcttggtatgtgcggccttttactgaagcatgggttactt  
accaggggctatttctttaaagaaaactggcctcttttcttggtagttattaattgccaata  
gttcttggttagaggtgggactctgtgcatacttctctctccatgttggaactttggect  
agcttaggcttgcaagtccttggtgatgctgttataaccaccatgaattcatatgtgaag  
ccaccctgttgatctggaagaccctgttctcttgagccacccaccacctttagccctta  
cactcttcagccctcttctcaatgatgcctatgccttggaagaagggtataggatatgaa  
ctcttagacacattttataatttgtttctccatttcacatataacatcatatcacattg  
ttgcataatatgtaatcatttttagaaggaaggagaggacttattgctcataaatgtccttt  
atgttgctgtgattggaccctcactatttaatttgtaattatttgaccacaagtgaggc  
atttcatatttcccaaatgtgaaagaggtaaaaaatctttgggttatgaaatggggatta  
catttttttctggctacaacatttacctcttggaagggggaatttaactaggtcatagct  
tgaactatgc

Fig. 16

## INTRON2/EXON3/INTRON3/EXON4/INTRON4

accacaaaatttttccttttagaaaccctgctggattttacaaaggcctggagctaatttag  
atgaattgtgattaccttaaaaggaaaggttgacacaggcctcaaatcaaacgtctctccc  
tgccatttcgggcctttgaccttctgcgagttggtcaccaggagatccttagctagtaaca  
aattactattggcctaagccaagctcataattaggtaaagaacactgcatttcaagttcaa  
attccttctattatatccagtgtatttagccttctcgtcagcgtggaataatcacgtgtta  
actgccagtcctttagaaacgatgcctttgttgaaaatgcttttataaaccacaaaaatt  
tagtctaaatgaagtttggtttgtaaatgggtccactcttggaataatgtatgtctttcc  
tttggtggtgacctcagcttcactagaaacttcttggtctatttctagcctgaaaggaat  
gcagagacaattctgttgcttctgatcatccgctgaccacacacctgcttatgaatgtac  
tttatataagatgatttagtttatgtgtttgtgtctgttggaatgaaaacaatggaaagagt  
tatatgtatctactaatctgagaggaaatagtgaaggcaagagatggtaaagtgtttggaa  
agatcaagaaatgcgccaaggaagacctaaaggtagttttgtaccactgctgtgctttg  
tagtttgagaggtcttcccttgagagcaaaagagctactcacaacatggctgtgcaagaat  
ggcctgtctatttttaggtgagacctcaacagaaatttaggaaggtcttctgtacctacag  
atagaggtgggtgggagactgttctgtctttgagaggtgtacaggtgggcaactaattaca  
gagaacaagaacatgattttgtactggaaaagtctcgtctcaaagtaggcaatgactgaga  
atagctttggacagctctgttatctctaacttatggtctagttgacagaagctctgtgtgcca  
ggcttcagtgtttctcattaaatgctaaatcccaagcagttcccatctcaaaaaccttcca  
gatggaaatatctgaaaaacaactttctaaaaaatataataacatagaaaggaaaaa  
aaacccgagtagtctattgttcagttgtctcatagcaatactattttatgttaaagtga  
cagtttggtgctgtgtgcttaagcttgcctcaaagctttgtggggagtgttgattcagg  
cagtgctctttgtgagagcacagatgtcagggaagaacacactcctggcagccacctgcaa  
atgtgatctctttaaactaccaatatataagtaaaggtaacctctatagccttcat  
tcatgtcatagaggaaaccttgtcatggcgttagcactgacagtgaagaacactggggaag  
gaagtcaaatttacatgaaggatgagtttagtatggcttatgatgtagaaagtatacgtga  
tgccagccgtgcatttacatgtgtcacacagtcagtttttaatttctctgtagtatt  
tatattgtgaaatgatgataggcgataggttgttagttgtttgtttgtttgtttgtttgt  
tttttattaatgacctgtaggatctgtgcttttttttttttttttttttttttttttttt  
ttcctttgttactgaattcatgtgggtaattggcctgtacaccttagaggttcgggatgt  
atgatgggtagcagtcagaagggtcagggctgtgagcaagagcttgaaggactaatacta  
ccttgggagagctgagatgacagctcattagctcttcttgcagggaaatgcagccaagca  
aaggaagctcttctggatgtgttttgccttcttctcttatgttttcaaatatttgagcaag  
ttgcacaaatgttacctctctgcaccttctcacatcttataggtctttttattttttaa  
gcataaagaggcaataaataatgggttttggtctttttattttccagACAGCACCTGGAGT  
ACATAAAAGATTTTCCGGAAAGTAAAAATCTCATCTCAGCGTTTCAGAAGCCGGATCAA  
GGCATCGTGACGCCTCTGCAGTATCCAGTTGAAAAGTCTCTGGCAGGGGGCCACAAGCTC  
CCACAGgtgtgctgcccagttttgaaagtattgcagagtttcgggttggggggcggggcat  
gtgggttaaactattcataccacgtcataaacaggaagacttataagaaaaacgcaacaa  
ggagagagctggagccagccaatcaggtgagaggtaatgacacaaaggggccataagatgc  
tctaacacttacctttttcataaaaacgtagaactatggaaagtatgattccgggatatt  
taatctctttatttctagcttattgtccctcttctcttcttcttcttcttcttcttctt  
gcttcagtaaccacgcagtgataatttgaaagatggcacaggaaggctagagctagcttg  
gttctgtgtactctgactcctggactgatgaccttgaatgaatgagtgaatttttactg  
tagcaaaggaattcaaattagcaaacttggtgttttaccagcatgcatgattctcctgag  
caaagtatttttagtgagttgttgagctcaggagtgaacacgcccagccagctgagctct  
tgttgttttttatgaaaaagggttagtattttgttttgagtgaagaatttgccatgatt  
gatgcagttaatatttatgataaaatgagttctttaaatttttgaggacaagaaccaggctt  
aggcataagttgataattagactcaaattggaattaaattaatgaatagcaaagcctttga  
aatgaaatatagatagaactctaatttagaatgcaactaactttcttctctgtccagtagt  
ctaattcttttgatgaatttttaataataatttttaaatagctgcatgaggcatagcattag  
gttcaatatatacaatttgagttaaactatttttaataactgaacaactttgtctctgatg

Fig. 17 (sheet 1 of 2)

tttgctattctttttcttttagGGAGAAGAGATTCTGATATCTGCCTGAATGCACCATGAag  
aagagtaactcactctgtactctattttcaataatttaagtatgtgccaagtcttgatatat  
tgtagatagtagacagttcagcgagctgcaaagtccttttaagcacgactctagtcctgtaat  
agcatctcgctgatgtcaaagctgcaagtggacgtggtagataacaagtgcctttaaagtg  
cggattggaaaactgaatcccttagggatatttttagttgccgtattttacaaatgagaaaca  
atttaaaagcacaatgaagactttacacaagattaatgtcgtttcttgccaaatgtaata  
aaataaaaatgatctcttttttttttttttgggttaaaagcggcatttttctgtgacttattt  
catagctattgatttgcaagcgtcttgcttgacttcatggtaccaaggtgggtgggtgca  
gatataatagctgtgggtgccaaggcttcttttgggatttatgaaggtgtgtgctcaaaga  
agtactagataaaagagaaaagactatttgtttaaatttgaaatccaccttcttgccctagag  
ttcttctgctaagacagttaaatccttggactagtggggtgtgggttaaggcatacttttcat  
tccagcactagagacatggaagtggggcgatcaggagtttagtatcatttggctacaggac  
acccttcttcaaaaacaaaataagcctgcagagcaaactggaatttacaagcctatatattt  
aaagtaggctttaaatgttcattgtgcttgatgctcaggttttgaaattatgtagaaacct  
gtgaaagatgcatgaattttgggtttcaaatatctgggcccccaatcgttccct

## mSAP Intron/exon sequences

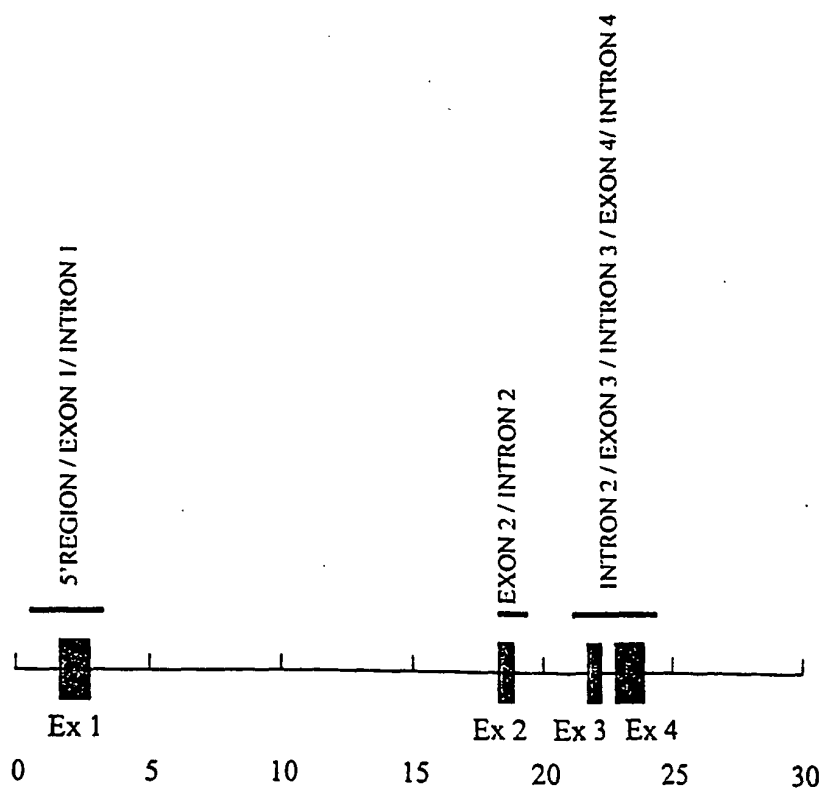
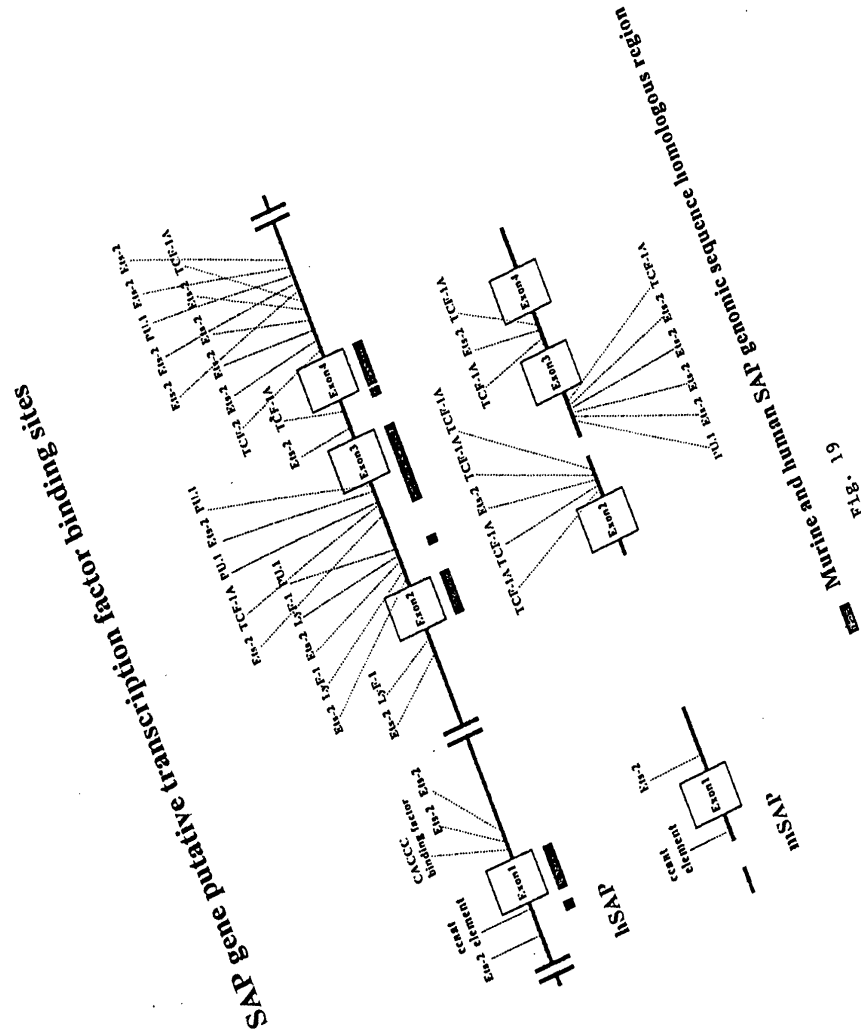


Fig. 18



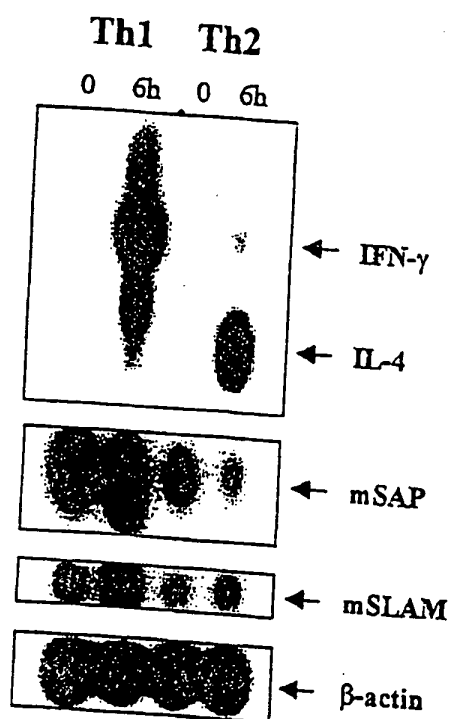


Figure 20

## COS CELLS TRANSFECTED

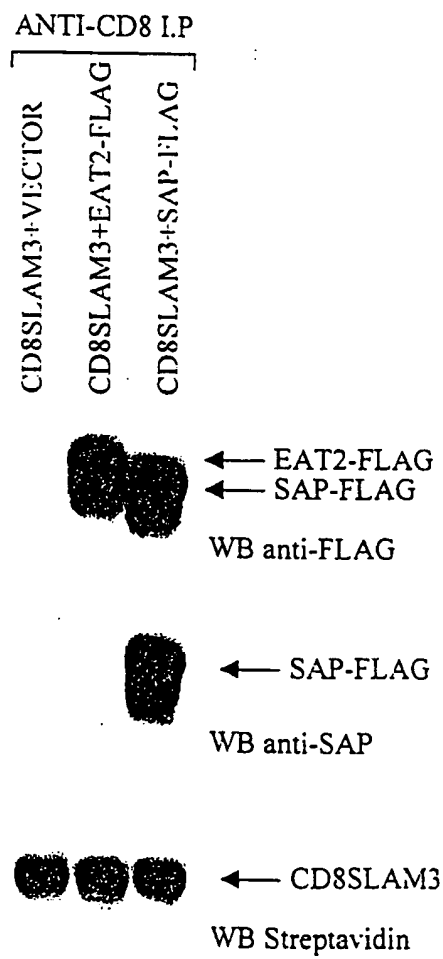


Fig. 21

## SEQUENCE LISTING

&lt;110&gt; Beth Isra l Deaconess Medical Center

<120> METHODS AND REAGENTS FOR THE UTILIZATION  
OF SAP FAMILY MEMBER PROTEINS, NOVEL SIGNAL TRANSDUCTION  
REGULATORS

&lt;130&gt; 01948/061WO2

&lt;140&gt; PCT/US98/xxxxx

&lt;141&gt; 1998-11-19

&lt;150&gt; 08/976,096

&lt;151&gt; 1997-11-21

&lt;150&gt; 60/099,160

&lt;151&gt; 1998-09-04

&lt;160&gt; 31

&lt;170&gt; FastSEQ for Windows Version 3.0

&lt;210&gt; 1

&lt;211&gt; 231

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 1

cagttgagaa gaagaggtaa aacgaacccat taccagacaa cagtggaaaa aaaaagcctt	60
acgatctatg cccaagtcca gaaaccaggt cctcttcaga agaaacttga ctccttccca	120
gctcaggacc cttgcaccac catatatgtt gctgccacag agcctgtccc agagtctgtc	180
caggaaacaa attccatcac agtctatgct agtgtgacac ttccagagag c	231

&lt;210&gt; 2

&lt;211&gt; 77

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 2

Gln	Leu	Arg	Arg	Arg	Gly	Lys	Thr	Asn	His	Tyr	Gln	Thr	Thr	Val	Glu
1				5					10					15	
Lys	Lys	Ser	Leu	Thr	Ile	Tyr	Ala	Gln	Val	Gln	Lys	Pro	Gly	Pro	Leu
			20					25					30		
Gln	Lys	Lys	Leu	Asp	Ser	Phe	Pro	Ala	Gln	Asp	Pro	Cys	Thr	Thr	Ile
		35				40					45				
Tyr	Val	Ala	Ala	Thr	Glu	Pro	Val	Pro	Glu	Ser	Val	Gln	Glu	Thr	Asn
	50					55					60				
Ser	Ile	Thr	Val	Tyr	Ala	Ser	Val	Thr	Leu	Pro	Glu	Ser			
65					70						75				

&lt;210&gt; 3

&lt;211&gt; 2226

&lt;212&gt; DNA



&lt;213&gt; Homo sapiens

&lt;400&gt; 3

```

ggttgacttg tgcctggctg cagtagcagc ggcattctcc ttgcacagtt ctctctctcg      60
gcctgcccac gagtccacca ggccatggac gcagtggctg tgtatcatgg caaaatcagc      120
agggaaaacc gcgagaagct cctgcttgcc actgggctgg atggcagcta tttgctgagg      180
gacagcgaga gcggtgccagg cgtgtactgc ctatgtgtgc tgtatcacgg ttacatttat      240
acataccgag tgtcccagac agaaacaggt tcttggagtg ctgagacagc acctggggta      300
cataaaagat atttccggaa aataaaaaat ctcatctcag catttcagaa gccagatcaa      360
ggcattgtaa tacctctgca gtatccagtt gagaagaagt cctcagctag aagtacacaa      420
ggtactacag ggataagaga agatcctgat gtctgctga aagcccatg aagaaaaata      480
aaacaccttg tactttatct tctataatct aaatatatgc taagtcttat atattgtaga      540
taatacagtt cgggtgagta caaatgcatt tctaaagcca ttgtagtctt gtaatggaag      600
catttagcat gtcgtcaaag ctgaaatgga cttttgtaca tagtgaggag ctttgaaacg      660
aggattggga aaaagtaatt ccgtaggtta ttttcagtta ttatatttac aaatgggaaa      720
caaaaggtaa tgaatacttt ataaaggatt aatgtcaatt cttgccaaat ataaataaaa      780
ataatcctca gtttttgtga aaagctccat ttttagtgaa atattatctt atagctacta      840
attttaaaat gtcttgcttg attgtatggt ggggaagtgg ctggtgtccc ttgtctttgc      900
caagtctctc actagctatg gtgtcatagg ctcttttggg atttttgaag ctgtatactg      960
tgtgtataaa caagcactaa acaaagagtg aaggatttat gtttaattct gaaagcaacc     1020
ttcttgccca gtgttctgat attggacagt aaaatccaca gaccaacctg gagtggaaaa     1080
tcttataatt taaaatatgc tctaaacatg tttatcgtat ttgatgctac aggatttgaa     1140
attgtattac aaatccaatg aaatgagttt ttcttttcat ttacctctgc cccagttggt     1200
tctactacat ggaagacctc attttgaagg gaaatttcag cagctgcagc tcatgagtaa     1260
ctgatttgta acaagcctcc ttttaaagta accctacaaa accactggaa agtttatggt     1320
tgtattatct ttaaaaaaaa ttccaagtga ttgaaaccta cagagatac agaattttat     1380
gcggcatttt cttctcacat ttatattttt gtgattttgt gattgattat atgtcacttt     1440
gctacagggc tcacagaatt cattcactca acaaacataa tagggcgctg agggcataga     1500
agtaaaaaaca cctggctcct gctctcagtt cactgtcttg ttggacgaga aaacaataac     1560
gataaaaagac agtgaaagaa aataacgata aaagacagtg aaagaaaata acaataaaaag     1620
acaagggaaa aataacaatg aaagttgata agtacatgat aagcgagggt ccccggtgtg     1680
aggtagatct ggtctttaga ggcagataga taggtcagtg caaatactct ggtccatggg     1740
ccatatgaaa aggctaagct tcactgtaaa ataataactg ggaattctgg attgtgtatg     1800
ggtgttggtg aacttggttt taattagtga actgctgaga gacagagcta ttctccatgt     1860
actggcaaga cctgatttct gagcatttaa tatggatgcc gtgggagtac aaaagtggag     1920
tgtggcctga gtaatgcatt atgggtggtt taccatttct tgaggtaaaa gcatcacatg     1980
aacttgtaaa ggaattttaa aatcctactt tcataataag ttgcatagggt ttaataatct     2040
ttaattatat ggcttgagtt taaattgtaa taggcgtaac taattttaac tctataatgt     2100
gttcattctg gaataatcct aaacatatga attatgtttg catgttcact tccaagagcc     2160
tttttttgaa aaaaagcttt ttttgaatca tcaagctttt cacatttaaa taaagtgttt     2220
gaaagc

```

&lt;210&gt; 4

&lt;211&gt; 128

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 4

```

Met Asp Ala Val Ala Val Tyr His Gly Lys Ile Ser Arg Glu Thr Gly
 1             5             10             15
Glu Lys Leu Leu Leu Ala Thr Gly Leu Asp Gly Ser Tyr Leu Leu Arg
          20          25          30
Asp Ser Glu Ser Val Pro Gly Val Tyr Cys Leu Cys Val Leu Tyr His
          35          40          45
Gly Tyr Ile Tyr Thr Tyr Arg Val Ser Gln Thr Glu Thr Gly Ser Trp

```

50	55	60
Ser Ala Glu Thr Ala Pro Gly Val His Lys Arg Tyr Phe Arg Lys Ile		
65	70	75
Lys Asn Leu Ile Ser Ala Phe Gln Lys Pro Asp Gln Gly Ile Val Ile		80
	85	90
Pro Leu Gln Tyr Pro Val Glu Lys Lys Ser Ser Ala Arg Ser Thr Gln		95
	100	105
Gly Thr Thr Gly Ile Arg Glu Asp Pro Asp Val Cys Leu Lys Ala Pro		110
	115	120
		125

<210> 5  
 <211> 381  
 <212> DNA  
 <213> Mus musculus

<400> 5	
atggatgcag tgactgtgta ccacggcaaa atcagcaggg agaccgggga gaagctctta	60
ctcgctaccg ggctggatgg aagctatctg ctgcgagaca gcgagagtgt ccctggcgtg	120
tactgcctgt gtgttttgta tcaagggttac atctacacat atcgagtgtc ccagacagaa	180
acaggttctt ggagtgccga gacagcacct ggagtacata aaagattttt ccggaagta	240
aaaaatctca tctcagcgtt tcagaagccg gatcaaggca tcgtgacgcc tctgcagtat	300
ccagttgaaa agtcctctgg caggggccca caagctccca cagggagaag agattctgat	360
atctgcctga atgcaccatg a	381

<210> 6  
 <211> 126  
 <212> PRT  
 <213> Mus musculus

<400> 6	
Met Asp Ala Val Thr Val Tyr His Gly Lys Ile Ser Arg Glu Thr Gly	
1	5
Glu Lys Leu Leu Leu Ala Thr Gly Leu Asp Gly Ser Tyr Leu Leu Arg	10
	15
	20
Asp Ser Glu Ser Val Pro Gly Val Tyr Cys Leu Cys Val Leu Tyr Gln	25
	30
	35
Gly Tyr Ile Tyr Thr Tyr Arg Val Ser Gln Thr Glu Thr Gly Ser Trp	40
	45
	50
Ser Ala Glu Thr Ala Pro Gly Val His Lys Arg Phe Phe Arg Lys Val	55
65	70
Lys Asn Leu Ile Ser Ala Phe Gln Lys Pro Asp Gln Gly Ile Val Thr	75
	80
	85
Pro Leu Gln Tyr Pro Val Glu Lys Ser Ser Gly Arg Gly Pro Gln Ala	90
	95
	100
Pro Thr Gly Arg Arg Asp Ser Asp Ile Cys Leu Asn Ala Pro	105
	110
	115
	120
	125

<210> 7  
 <211> 332  
 <212> DNA  
 <213> Homo sapiens

<400> 7	
atggacgcag tggctgtgta tcatggcaaa atcagcaggg aaaccggcga gaagctcctg	60
cttgccactg ggctggatgg cagctatttg ctgagggaca gcgagagcgt gccaggcgtg	120

```
tactgcctat gtgtgctgta tcacggttac atttatacat accgagtgtc ccagacagaa 180
acaggttctt ggagtgtgta gcatttcaga agccagatca aggcattgta atacctctgc 240
agtatccagt tgagaagaag tcctcagcta gaagtacaca aggtactaca gggataagag 300
aagatcctga tgtctgcctg aaagcccat ga 332
```

<210> 8  
 <211> 76  
 <212> PRT  
 <213> Homo sapiens

```
<400> 8
Met Asp Ala Val Ala Val Tyr His Gly Lys Ile Ser Arg Glu Thr Gly
 1             5             10             15
Glu Lys Leu Leu Leu Ala Thr Gly Leu Asp Gly Ser Tyr Leu Leu Arg
             20             25             30
Asp Ser Glu Ser Val Pro Gly Val Tyr Cys Leu Cys Val Leu Tyr His
             35             40             45
Gly Tyr Ile Tyr Thr Tyr Arg Val Ser Gln Thr Glu Thr Gly Ser Trp
             50             55             60
Ser Ala Glu His Phe Arg Ser Gln Ile Lys Ala Leu
65             70             75
```

<210> 9  
 <211> 323  
 <212> DNA  
 <213> Homo sapiens

```
<400> 9
atggacgcag tggctgtgta tcatggcaaa atcagcaggg aaaccggcga gaagctcctg 60
cttgccactg ggctggatgg cagctatttg ctgagggaca gcgagagcgt gccaggcgtg 120
tactgcctat gtgtgctaca gcacctgggg tacataaaag atatttccgg aaaataaaaa 180
atctcatttc agcatttcag aagccagatc aaggcattgt aatacctctg cagtatccag 240
ttgagaagaa gtcctcagct agaagtacac aagggtactac agggataaga gaagatcctg 300
atgtctgcct gaaagcccca tga 323
```

<210> 10  
 <211> 58  
 <212> PRT  
 <213> Homo sapiens

```
<400> 10
Met Asp Ala Val Ala Val Tyr His Gly Lys Ile Ser Arg Glu Thr Gly
 1             5             10             15
Glu Lys Leu Leu Leu Ala Thr Gly Leu Asp Gly Ser Tyr Leu Leu Arg
             20             25             30
Asp Ser Glu Ser Val Pro Gly Val Tyr Cys Leu Cys Val Leu Gln His
             35             40             45
Leu Gly Tyr Ile Lys Asp Ile Ser Gly Lys
50             55
```

<210> 11  
 <211> 268  
 <212> DNA  
 <213> Homo sapiens

<400> 11  
 atggacgcag tggctgtgta tcatggcaaa atcagcaggg aaaccggcga gaagctcctg 60  
 cttgccactg ggctggatgg cagctatttg ctgagggaca gcgagagcgt gccagggcgtg 120  
 tactgcctat gtgtgctcat ttcagaagcc agatcaaggc attgtaatac ctctgcagta 180  
 tccagttgag aagaagtcc cagctagaag tacacaaggc actacaggga taagagaaga 240  
 tcctgatgtc tgcctgaaag ccccatga 268

<210> 12  
 <211> 62  
 <212> PRT  
 <213> Homo sapiens

<400> 12  
 Met Asp Ala Val Ala Val Tyr His Gly Lys Ile Ser Arg Glu Thr Gly  
 1 5 10 15  
 Glu Lys Leu Leu Leu Ala Thr Gly Leu Asp Gly Ser Tyr Leu Leu Arg  
 20 25 30  
 Asp Ser Glu Ser Val Pro Gly Val Tyr Cys Leu Cys Val Leu Ile Ser  
 35 40 45  
 Glu Ala Arg Ser Arg His Cys Asn Thr Ser Ala Val Ser Ser  
 50 55 60

<210> 13  
 <211> 443  
 <212> DNA  
 <213> Homo sapiens

<400> 13  
 ggaaactgtg gttgggcaga tacaatatgg agacaagtta gattcactat gctttacttc 60  
 ctatgaatgc aatgacacca tatacgtgtg tcctagtata tgtgacattt atatgtaata 120  
 ttaagctcaa atttaaagta tccattgttc ttttggaatc tttcagtaat ggaagtttat 180  
 tctttcagag gtatcacggt tacattttata cataccgagt gtcccagaca gaaacagggt 240  
 cttggagtgc tgagggtatag ttgtattttat ttttgcttct gggggtgtca aggagggtatt 300  
 tgaaatttag gctgggtttta taaaagagca aattatacat tattaagtat tcataagggt 360  
 taaatctcta aagctccaat ccaaattgt tcatggatca ttaagaaagc tttagagaat 420  
 tttggtccca gtctgtttta gcc 443

<210> 14  
 <211> 4011  
 <212> DNA  
 <213> Mus musculus

<400> 14  
 taagatattt tttctacttt tgtaaaggaa gtaaacttaa gtgccaagag tatgtgtaaa 60  
 ctcccacatc ataacatggc cacaggagaa ttcacacatt ttagagaagc tggtagaaga 120  
 acaaaggtag ttttgaaagg ttagagaaca agattctgaa atgtcctatg tactatacaa 180  
 aactgaccac ccttggcatc tttcttttaa tctacttggg ctccagaata gcctagggca 240  
 tttggaatac agacaattag cacctcagac agggatatcca cgtgctaaat cctgtcactg 300  
 ttcagagatc tcaggaggag acaagtttag acaagaagac cctttaaac tgcctgaca 360  
 ggttttctag ctttcaatgt tttggtgttg gacttgaatg gcatctatat tcagaacggt 420  
 ttcttttaaaa tataaggcac tcttctctta atgatagttt ttatttctga ggttttttaa 480  
 aaaacaatga gacaaatgcc tcctcctggt gagacttcag gatcttgctt ctctctcaaa 540  
 aaatgagcta cccctgtggt tttctccaaa gaagcttgct tgcttgcttg cttgcttgct 600  
 tcattcattc attcattcat ttattcattc acttatagca cagaatgttg aagctgaaaa 660  
 tgactactca ggggtcacct ggctcaattc catttgacag atggagaaaa taaggctcag 720

agagaggaaa	ccatttagtg	ataaaagggtg	gagtacaatt	tggacctcaa	aataccttcc	780
aattctcatc	gctgtttcta	agggagccaa	agagacacgc	actctctctc	tctctctctc	840
tctctctctc	tttctctctc	tctctctctg	tgtgtgtgtg	tgtgtgtcag	agagagagag	900
agacagagat	agacagagac	agtcagagac	cgacagagag	acagacagat	agacattgag	960
agtcagagaa	agagacagac	agacaagggg	gagagataat	ttcatattgt	tttcttgaca	1020
attttaattt	cagacattta	ttcattcaact	aaacccatcc	actctgtgcc	agtctctgag	1080
gatacagtgg	caaacaacat	atataagacc	ctgttgatat	gcaccttaag	gtcaagtgat	1140
agagacaggc	aacatacaag	tagattcttc	ctctcataaa	tggggcccat	tccttagctc	1200
tttgtctgtt	ggtgtcactg	ttatctcccc	ttctctgtta	tgtgatcttt	gtcttctctg	1260
actgtgggta	cagggacagg	gaaggagaa	agctatttca	gttctcttat	tctctacact	1320
tcaagcttag	tgctgcata	gcagcagagc	aagggtgtca	gtatttttca	tcaacactgc	1380
actggtgggc	tggctcacct	gtcagtttgc	ttgtgagatg	acaataagg	agtggatgtg	1440
agagttgtac	tattaccgta	taatttaact	gttctcttgt	gatgtgattt	tcgttgtgct	1500
gaagcccaact	gtacctggct	gctttgacaa	gcgtgctatt	ttagcaatgc	agaatgaaac	1560
agaaagaaaa	gaatgcatgg	gatgcttgct	ccgtcagatc	caggtttctc	tgtgcctgtg	1620
tttgtagatg	atgtgttcag	ggttgttcag	agacaaaaaa	aatgagggtta	ggttttgttt	1680
ttaacctatg	gaacactgaa	tgttctagca	gttactctct	aggtgggcta	ctgtccgtat	1740
gccttaataa	aagaagatta	cccccttctt	tcttttcata	tttaaaatgc	catcattata	1800
actgatttct	aaatagagtc	ctggttaggc	taaactctga	tgttttaaca	ttgatgtagg	1860
taggcattta	tgtatgagaa	tagaagtacc	acacatttat	caacacctgt	gccatgatgt	1920
aaaaatggat	tttatatccc	aatgattctg	gcctagcccc	atatgccaaa	gggtatatag	1980
aagttcacac	aggacaccaa	tggacagacc	aaactggaca	gaaaaagccc	acaaaccagc	2040
agctctacag	gaagaactac	agacagctga	gtaaaaatcg	aagtagaaga	agtggtcacc	2100
cgtgggaaga	gcacactaat	tggctgtcta	ctaccaaatg	gctatcccctg	aaaacataca	2160
tacaaatagc	accatacagg	cacaactgga	tgtgttttag	aattacacgt	atatacatat	2220
atgcatgcaa	taatacaata	atacaatagc	aatgttgttg	aagagcaggg	agggattcat	2280
gggaggggtt	caagggtgaa	aagggaaggg	agaaatgctg	tgattatatt	gtgattttaa	2340
taataataaa	aacaaagaag	taccaacaaa	agaatttagt	cttcttaagc	tccttagcta	2400
gatccactta	cacttttttt	tttttaatat	tttttcatag	ttttggattt		2460
tggacaaaac	attaaggtaa	aaatacttgt	ttgggctaag	tgtattcaag	gatttggcat	2520
atggcttcgg	gatagaagac	ctcatctcgc	aggccccgaa	ttccatcatc	agcattgaag	2580
aaaagaaaagc	tgtttcagga	ccatctagtc	ccaaagtctg	tagttcagta	gcccttgggtg	2640
atgctgtgga	agccttgacg	ccatgttctg	ctgtggtttt	gtagcttgac	tgtttttatg	2700
cttttatattt	gtcagtctgc	ttctgcccc	catgatgaac	acgtcccatt	cacttctatc	2760
accaattctt	ttgttcactc	gttaagacaa	gtcattgtca	atggcttcga	agtttgcagg	2820
aactacacaa	gaacaaaggc	actagcaagt	ttctcatctt	tgtgcaagct	ctgtttgcgt	2880
tgctggcaca	gctcaaatac	tgcaagtgtt	caccttttca	gcgtgcccc	ttccccacca	2940
caaacctttg	ttggagaata	gatgatcaaa	cccagacaa	tttgaagctg	ttaaactgct	3000
gagctggcct	gaattgaagt	gtgagaagag	gttttaaggc	aggtagacaa	catcctgttg	3060
tgaggggggtt	ctctctctct	ctctctctct	ctctctctct	ctctctctct	ctctctctct	3120
ctctctctct	ctctcaatac	agcccagtaa	gccactgcag	cggaacagc	agcagcaag	3180
tttccctttt	accatccctc	ttctggggcc	atcaaagagt	ttgctaggca	tggtatgcag	3240
gactgtgtac	cacggcaaaa	tcagcaggga	gaccggggag	aagctcttac	tcgtaccgg	3300
gctggatgga	agctatctgc	tgcgagacag	cgagagtgtc	cctggcgtgt	actgacctgtg	3360
tgttttgtga	gtatatctcg	tggtagtggc	aggttgttag	catgggtagc	ctgccctttc	3420
aggggagggc	tgggaggagt	ggcaggggat	gacagctttt	ggcagaggta	cctagggaa	3480
acaggcagca	ggaactgagt	actcttcag	ctctgacgta	tgggtcccag	gtcagccca	3540
gggatgtggt	gggcagtact	tgtacgtacg	ttccctcactg	cggattttat	caactgcatt	3600
tcaccagtgc	gtttgtggca	acttcactga	atacccttca	gaatgggaaa	acatggccta	3660
gaacttacct	ttactttgag	aaagacccat	actctggact	tcggctctga	ttccaagtag	3720
acaccatgat	ggtggagccc	cggaagttg	gactatttgt	attctggggc	atgaagtaag	3780
gcaaagggaag	gtctttctct	atcattctct	acaaatgtga	ctgaaaatgg	tacctaaaa	3840
gaaaagcacc	tagcaaatgc	ttgaagagtg	gttagccagg	aaatggaaatc	cattgtcttc	3900
agaatgcagc	ccagggaagg	cagaggtaga	cagatgtgtg	tgagtttcag	gtgactaacc	3960
tgggtttaca	tggcccggtc	atccaactag	gctacataga	taaaaaccaa	g	4011

<210> 15  
 <211> 2877  
 <212> DNA  
 <213> Mus musculus

<400> 15

gaaacccttt	ggagaccaa	cagggtcttg	gagtgccgag	gtacagctct	tttatttgtt	60
tttactttctc	tgggtataag	aaaagtgcct	gaaatctagg	ttgggtttaa	aaagagcagt	120
catactatac	actacaaaa	tatttcaatc	actaatgctc	aattctatag	tcattgatgg	180
ttcattttaga	agccttttaga	aaattgtggt	tccagactta	tttagccacc	attatgtgag	240
gtaggatcat	tttgggaaaa	tgaaaaaatt	tgaaggacca	tgtgttaact	atagtgtctt	300
tacaaacttt	tagaaagcat	ataacaataa	taacaacaac	aacaacaaca	acaacaacaa	360
caataataat	acaaaccta	aaaactaaaa	aaaaaaaaaa	agggaaacca	tatactaaat	420
gatactgatg	cacagattgg	ttctgtttgt	gataaagcaa	actgggattt	ctatttccta	480
ctagtactag	tacccttttg	gaactctaac	aggggattat	actacagcgt	gaaatcctgg	540
gccatgtcaa	atgcattcag	actccatgtg	tagttcttcc	tatagtcact	tttgatggaa	600
gcaatttgta	aatgggtct	acaaacattt	atcaagagga	cccatttcag	ggagttataa	660
ggcaatgcaa	ctaaagctct	tagctggaga	gtctgacata	cagtgcagcat	ccaaaaatgt	720
gaactactga	aaacttctca	gcagtatgat	tgtgtgtcat	ctggtgtaaa	tttaccaggc	780
acaagaattt	gaacagaggc	attttcaaac	atccaggcga	ttgctgtctg	ctgtgtgtgca	840
taggcaaggt	tttcaaagga	gccataaaaag	agcctctgga	tagtcggaaa	agagaagagg	900
gaaaattgta	gtttttat	aataaaaatc	ttctttgaga	atttcaaata	tatatacaca	960
atgaattcct	atctgctatt	gtagtgtatt	aaaaggatgt	tttatagaat	gatgttctcc	1020
tccatcttgg	gactccctag	gagtcacaaga	agaagaagag	gaggaggtga	gtgggtatac	1080
ttctgggtac	caatcttggc	ttcactgggt	tagatccctt	tagttttcct	tatgaactag	1140
ggctttgaat	aacattaatt	cagaatgaag	gtttcttatg	gctttgaaat	atttgtacac	1200
cactgtgaca	gtccagaaat	tattgaacta	tttctagaga	gcagggtgga	tagagcaatg	1260
catttatgaa	tcaattcatg	attctacagc	tgtgggtttc	aaactttagg	ctttagttgg	1320
tttctgacct	tatactagtt	tatcctgata	gtggaaatgc	ttcacaatga	tcttttgaaa	1380
gcatagatgt	caatatcaat	ttctattagt	aactaaataa	atctgtgcat	ttaaagaata	1440
ataaagtcta	gacaaaagtt	ttaagaatgt	cttacaatt	tgtcagtaaa	aagcaaagta	1500
taaatttcaa	gtatttgaat	tatcaaaaca	aattagaata	tattcaagta	tactgttate	1560
ttattaatgt	gttaatat	tttttaagtt	ttatttttaa	ctttcaggtt	cgtaaaaaat	1620
agatgaagga	tgaaaccag	ggtctcatgc	attttagaca	attgttatac	cattgaataa	1680
tattaccaaa	taattttcct	ttttattgcc	atatattcgc	tgtacatgtt	aatgcacttc	1740
ataagacaag	tttataatat	cttttgatta	tgttccctca	taaaatacta	tatccctttg	1800
tggccatgtt	gttattttgt	gtcaggatac	tatccaaggg	aaaatcaata	ccatttctaa	1860
taaaaactat	acttctttaa	caaactctatc	acttttaata	aataagcaat	ttgacttate	1920
atgaataata	attatgtata	gctgatacat	atgaatttgt	gcatgggttg	ctatatttaa	1980
tctttttcca	ccccagaaa	tagccttcag	atgagcaata	gaaaatttaa	tggagagcct	2040
ttgatTTTTT	TTTTTactt	aaaaaattag	ttctttgaga	attttgtaca	atatgttttg	2100
atcatattta	cctttgtctt	ctaactcctt	cagatccacc	cttccctctc	tattcacccg	2160
cctttttatac	tcctgtttat	ccctaagtct	gtcaatgcag	tttgtgtctac	ccaaatatte	2220
ttggatgtgc	ggccttttac	tgaagcatgg	tttacttacc	aggggctatt	ctcttaaaga	2280
aaactggctc	ttttcctggg	agttattaat	tgccaatagt	tccttggtta	gaggtgggac	2340
tctgtgcata	cttcctctct	ccatgttgga	ctttggccta	gcttaggctt	gcacaagtct	2400
tgtgtatgct	gttataacca	ccatgaattc	atatgtgaag	ccaccctgtt	gtatctggaa	2460
gacctgtgtt	ccttggagcc	accaccacc	tttagccctt	acactcttca	gccctcttcc	2520
tcaatgatgc	ctatgccttg	ggaagaaggg	ataggatatg	aactcttaga	cacattttat	2580
aatttgTTTT	ctcccatctc	acatataaca	tcatatcaca	ttgttgcata	tatgtaatca	2640
TTTTtagaag	gaaggagagg	acttattgct	cataaatgtc	ctttatgttg	tctgattgga	2700
ccctcactat	ttaatatgtt	aattatttgc	accacaagtg	caggcatttc	atatttcccc	2760
aatgtgaaa	gaggtaaaaa	atctttgggt	tatgaaatgg	ggattacatt	tttttctggc	2820
tacaacattt	acctcttgga	aaggggaaat	ttaactaggt	catagcttga	actatgc	2877

<210> 16  
 <211> 4018  
 <212> DNA  
 <213> Mus musculus

<400> 16

acaaaaaat	ttttccttta	gaaaccctgc	tggattttac	aaaggcctgg	agctaattta	60
gatgaattgt	gattacctta	aaaggaaagg	ttgcacaggg	cttcaaataca	aacgtctctc	120
cctgccattt	cgggcctttg	accttctgcg	agttggctac	caggagatcc	ttagctagta	180
acaaattact	attggcctaa	gccaagctca	taattaggta	aagaacactg	catttcaagt	240
tcaaattcct	tctattatat	ccagtgcctat	tagccttctc	gtcagcgtgg	aataatcacg	300
tgttaactgc	cagtcctctgt	agaaacgatg	cctttgttga	aaatgctttt	ataaaccaca	360
aaaatttagt	ctaaatgaag	tttggtttgt	aaatgggtcc	actcctggaa	aatatgtatg	420
tctttccttt	ggctgggtgac	ctcagcttca	ctagaaactt	cttggctctat	tcctagcctg	480
aaaggaatgc	agagacaatt	ctgttgcttc	tgatcatccg	cgtgaccaca	cacctgctta	540
tgaatgtact	ttatataaga	tgattagttt	atgtgtttgt	gtctgtggaa	tgaaaacaat	600
ggaaagagtg	tatatgtatc	tactaatctg	agaggaaata	gtgaaggcaa	gagatggtaa	660
agtgtttgga	aagatcaaga	aatgcgccaa	ggaagaccta	aggtagtatt	ttgtacccac	720
tgcgtgcctt	tgtagtttga	gaggtcttcc	ttggagcaaa	agagctactc	acaaacatgg	780
ctgctgcaag	aatggccctg	ctatttttag	gtgagacctc	aacagaaatt	taggaagggtc	840
ttctgatcct	acagatagag	gtgggtggga	gactgtttct	gtccttgaga	gtgtacaggt	900
gggcactaat	ttacagagaa	caagaacatg	atttgtactg	gaaaagtcct	cgcttcaaa	960
taggcaatga	ctgagaatag	ccttgacacg	tctgttatct	ctaacttatg	gtctagttga	1020
cagaagtctg	tgtgccaggc	ttcatgtttt	ctcattaaat	gctaaatccc	aagcagttcc	1080
catctcaaaa	accttccaga	tggaatatct	gaaaaacaac	tttctaaaaa	atataataac	1140
atagaaagga	aaaaaaaaaa	aacccgagta	gtcattgttc	agttgctcat	agcaatacta	1200
ttttatttta	tgtaaaatga	cagtttgccg	ctgggtgccta	aagcctgcct	caaaagcttt	1260
gtggggagtg	ttgatttcag	gcagtgctct	tgtgagagca	cagatgtcag	ggaaagaaca	1320
cactcctggc	agccacctgc	aaatgtgatc	tctttaaaaa	ctaccaatat	ataagtaaag	1380
gtaaccttct	atagccttca	ttttcatgtc	atagaggaaa	ccttgctcatg	gcgtagcact	1440
gacagtgaga	acaactgggg	taaggaagtc	aaatttcatg	gaaggatgag	ttagtatggc	1500
ttatgatgta	gaaaagtata	cgtgatggcc	agccgtgcat	ttacatgtgt	cacaacagtc	1560
agtttttaat	tttctctgta	gtattttata	ttgtgaaatg	atgataggcg	ataggttggt	1620
agttgtgtgt	gttttggttg	ctcaggcttt	ttattaatga	cctgtaggat	ctgtgctttt	1680
tttttttttt	tttgatgttt	aactgcattt	cctttgttac	tgaattcatt	gtgggtaatg	1740
gcctgtacac	ccttagaggt	tccggatgta	tgatgggtag	cagtcaagaa	gggcagggct	1800
gtgagcaaga	gcttgtaagg	actaatacta	cccttgggag	agctgagatg	acagctcatt	1860
agctcttctt	gcagggaagt	gcagccaagc	aaaggaaagt	ccttctggatg	ttgttttgct	1920
ttcttcctta	tgttttcaaa	tatttgagca	agttgcacaa	atgttacctc	tctgcatcct	1980
tctcacatct	tataggtctt	tttatttttt	taagcattaa	gaggcaataa	aataatgggt	2040
ttggctcttt	atttttccag	acagcacctg	gagtacataa	aagatttttc	cggaaagtaa	2100
aaaatctcat	ctcagcgttt	cagaagccgg	atcaaggcat	cgtgacgcct	ctgcagtatc	2160
cagttgaaaa	gtcctctggc	aggggcccac	aagctcccac	aggtgtgctg	cccagttttg	2220
aaagtattgc	agagtttcgg	gttggggggc	ggggcatgtg	ggtaaacctt	ttcataccca	2280
cgtcataaac	aggaagactt	ataagaaaaa	cgcaacaagg	agagagctgg	agccagccaa	2340
tcaggtgaga	ggtaatgaca	caaagggggc	ataagatgct	ctaacactta	cctttttcat	2400
aaaaccgtag	aactatggaa	agtatgattc	cgggatattt	taatctcttt	atttctagct	2460
tattgtccct	cttcctcttt	gtagacagta	aataccactt	tgcttcagta	acccacgcct	2520
gtacaatttg	aaagatggca	caggaaggct	agagctagct	tggttcctgt	gattctgact	2580
cctggactga	tgacctttga	atgaatgagt	gaaattttta	ctgtagcaaa	ggaattcaaa	2640
ttagcaaaact	tggctgtttt	accagcatgc	atgattctcc	tgagcaaaagt	tatttttagt	2700
gagtttggtga	gctcaggagt	gaacacgcca	gccagctgat	gtctctgttg	ttcttttatg	2760
aaaaaggggt	agtattttgt	tttgagttag	aagatttgcc	atgattgatg	cagtttaatat	2820
ttatgataaa	atgagttctt	aaaattttga	ggacaagaac	caggcttagg	cataagttga	2880
taattagact	caaatgtgaa	ttaaattaat	gaatagcaaa	gcctttgaaa	tgaaatatag	2940

```

atagaactct aatttagaat gcaactaact ttcctttctg tccagtagtc tattcttttg 3000
gatgaatttt aaataataat tttaaatagc tgcattgaggc atagcattag gttcaatatt 3060
acaaatttga gttaaactat tattttaata ctgaacaact ttgtcctgat gtttgctatt 3120
ctttttcttt agggagaaga gattctgata tctgcctgaa tgcaccatga agaagagtaa 3180
ctcactctgt actctatttt caataattta agtatgtgcc aagtcttgta tattgtagat 3240
agtacagttc agcgagctgc aaatgccttt taagcacgac tctagtcctg taatagcatc 3300
tcgcctgatg tcaaagctgc aagtggacgt ggtagataac aagtgcttta aagtgcggat 3360
tggaactctg aatcccttag ggtattttta gttgcccgtat ttacaaatga gaaacaattt 3420
aaaagcacaa tgaagacttt acacaagatt aatgtcgttt cttgccaaat gtaataaaaa 3480
taaaaatgat ctcttttttt tttttttggt taaaagcggc atttttcgtg acttatttca 3540
tagctattga tttgcaagcg tcttgcttga cttcatggta ccaagggtgg tgggtgtgcag 3600
atataatagc tgtggtgccc aaggcttctt ttgggattta tgaagggtgtg tgctcaaaga 3660
agtactagat aaagagaaaa gactatttgt ttaaatttga aatccacctt cttgcctaga 3720
gttcttctgc taagacagtt aaatcttttg actagtgggg tgtggttaagg catacttttc 3780
attccagcac tagagacatg gaagtggggc gatcaggagt ttagtatcat ttggctacag 3840
gacacccttc ttcaaaaaca aaataagcct gcagagcaaa ctggaattta caagcctata 3900
ttttaaaagta ggctttaaat gttcattgtg cttgatgctc aggttttgaa attatgtaga 3960
aacctgtgaa agatgcatga attttgggtt tcaaatatct gggccccaat cgttccct 4018

```

&lt;210&gt; 17

&lt;211&gt; 29

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 17

```
gcctggctgc agtagcagcg gcatctccc 29
```

&lt;210&gt; 18

&lt;211&gt; 28

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 18

```
atgtacaaaa gtccatttca gctttgac 28
```

&lt;210&gt; 19

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 19

```
gccctacgta gtgggtccac ataccaacag 30
```

&lt;210&gt; 20

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens

&lt;400&gt; 20

```
gcaggaggcc cagggaatga aatccccagc 30
```

&lt;210&gt; 21

&lt;211&gt; 30

&lt;212&gt; DNA

&lt;213&gt; Homo sapiens



<400> 21  
 ggaaactgtg gttgggcaga tacaatatgg 30  
  
 <210> 22  
 <211> 29  
 <212> DNA  
 <213> Homo sapiens  
  
 <400> 22  
 ggctaaacag gactgggacc aaaattctc 29  
  
 <210> 23  
 <211> 29  
 <212> DNA  
 <213> Homo sapiens  
  
 <400> 23  
 gctcctcttg cagggaaatt cagccaacc 29  
  
 <210> 24  
 <211> 31  
 <212> DNA  
 <213> Homo sapiens  
  
 <400> 24  
 gctacctctc atttgacttg ctggctacat c 31  
  
 <210> 25  
 <211> 31  
 <212> DNA  
 <213> Homo sapiens  
  
 <400> 25  
 gacagggacc taggctcagg cataaactga c 31  
  
 <210> 26  
 <211> 28  
 <212> DNA  
 <213> Homo sapiens  
  
 <400> 26  
 atgtacaaaa gtccatttca gctttgac 28  
  
 <210> 27  
 <211> 15  
 <212> PRT  
 <213> Homo sapiens  
  
 <400> 27  
 Cys Val Glu Lys Lys Ser Leu Thr Ile Tyr Ala Gln Val Gln Lys  
 1 5 10 15  
  
 <210> 28  
 <211> 15  
 <212> PRT

<213> Homo sapiens

<400> 28

Cys	Val	Glu	Lys	Lys	Ser	Leu	Thr	Ile	Tyr	Ala	Gln	Val	Gln	Lys
1				5					10				15	

<210> 29

<211> 15

<212> PRT

<213> Homo sapiens

<400> 29

Cys	Val	Glu	Lys	Lys	Ser	Leu	Thr	Ile	Phe	Ala	Gln	Val	Gln	Lys
1				5					10				15	

<210> 30

<211> 18

<212> PRT

<213> Homo sapiens

<400> 30

Cys	Thr	Thr	Ile	Tyr	Val	Ala	Ala	Thr	Glu	Pro	Val	Pro	Glu	Ser	Val
1				5					10				15		

Gln Glu

<210> 31

<211> 12

<212> PRT

<213> Homo sapiens

<400> 31

Cys	Thr	Val	Tyr	Ala	Ser	Val	Thr	Leu	Pro	Glu	Ser
1				5					10		

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US98/24976

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(6) : C07K 16/00, C07H 21/04; C12P 21/06; A61K 38/00

US CL : 530/388.75; 536/23.5; 435/69.1; 514/12

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 530/388.75; 536/23.5; 435/69.1; 514/12

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, MEDLINE, SCISEARCH, BIOSIS, EMBASE, BIOTECHDS, LIFESCI, NTIS, WPIDS

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Database Genbank, Accession No. AA255258, mz77h04.r1 soars mouse lumph node NbMLN Mus, 14 March 1997, see the entire document.	7
X,P	Database sptremb16, Accession No. O60880, Duncan's disease SH2-protein protein (DHSP), 01 August 1998, see the entire document.	15, 20
X,P	Database Genbank, Accession No. AL022718, Human DNA sequence from clone 1052M9 on vchromosome Xq25. Contains the SH2D1A gene for SH2 domain protein, Duncan's disease (lumphoproliferative syndrome) (DHSP), see the entire document.	7

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E* earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A* document member of the same patent family
*O* document referring to an oral disclosure, use, exhibition or other means	
*P* document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

01 FEBRUARY 1999

Date of mailing of the international search report

25 FEB 1999

Name and mailing address of the ISA/US  
Commissioner of Patents and Trademarks  
Box PCT  
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

MARYAM MONSHIPOURI

Telephone No. (703) 308-0196

Form PCT/ISA/210 (second sheet)(July 1992)\*

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US98/24976

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X,P	SAYOS et al. The X-linked lymphoproliferative disease gene product SAP regulates signals induces through the co-receptor SLAM. Nature. 01 October 1998, Vol. 395, pages 462-469, see the whole document specially figure 5.	1-21

Form PCT/ISA/210 (continuation of second sheet)(July 1992)\*

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US98/24976

**Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)**

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

Please See Extra Sheet.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: | - 2 |

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.  
☐ No protest accompanied the payment of additional search fees.

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US98/24976

### BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING

This ISA found multiple inventions as follows:

This international searching authority has found 5 inventions claimed in the international application covered by the claims indicated below:

Group I, claims 1-21, drawn to SAP polypeptides, its fusion proteins, fragments, the polynucleotides encoding the polypeptides and a method of treating disease involving administering SAP family polypeptides.

Group II, claims 22-26 and 31-33, drawn to a method of identifying compounds that modulate the SH2 domain of SAP polypeptides utilizing a SAP family member polypeptide-encoding gene.

Group III, claims 27-30, drawn to a method of identifying SAP polypeptide mimetics utilizing a SLAM polypeptide with a SAP polypeptide.

Group IV, claims 34-37, drawn to a method of increasing antigen-specific T cell activation in a mammal, comprising administering to a mammal a compound which increases the biological activity of SAP polypeptides.

Group V, claims 38-41, drawn to a transgenic mammal, having a knockout mutation in an endogenous SAP family member protein-encoding nucleic acid sequence.

The inventions listed as groups I-V do not relate to a single inventive concept because they are considered to be 5 different categories of invention and are not drawn to combinations of categories (i.e. categories 1-5), specified in 37 CFR section 1.475(b).